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(54) Thermal recording sheet

Wärmeempfindliches Aufzeichnungsblatt

Feuille pour l'enregistrement thermique

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(56) References cited:
EP-A- 0 615 859 EP-A- 0 620 122
CA-A- 1 237 145 GB-A- 2 079 480

• DATABASE WPI Week 8713, Derwent
Publications Ltd., London, GB; AN 87-091461 &
JP-A-62 042 883 (RICOH KK) 24 February 1987
• DATABASE WPI Week 8715, Derwent
Publications Ltd., London, GB; AN 87-104704 &
JP-A-62 051 480 (RICOH KK) 30 August 1985

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Description

This invention relates to a thermal recording sheet with improved stability of the ground color and to a thermal recording sheet having a reversible recording ability with a bright color tone.

In general, to obtain a thermal recording sheet, a colorless or pale colored electron-donating dye and a color developer such as phenolic compounds are individually pulverized to fine particles, mixed with each other, and a binder, a filler, a sensitizer, a slip agent, and other additives are added to obtain a coating color, which is coated on a substrate such as paper, synthetic paper, films, plastics, or the like. The coated sheet is color developed by an instantaneous chemical reaction by heating with a thermal head, a hot stamp, laser light, or the like to obtain a visible record.

The thermal recording sheet is used in a wide variety of applications such as measuring recorders, a terminal printer for computers, facsimile, automatic ticket vendors, bar-code labels, and the like. However, with recent diversification and advance in recording apparatus for thermal recording sheet, the thermal recording sheet is required to meet increased requirements. For example, with increasing speed of recording, it is required to obtain a high-density and sharp color image even with a smaller thermal energy, whereas, in view of storage stability of the recording sheet, it is required to have improved light resistance, oil resistance, water resistance, and solvent resistance.

On the other hand, with increasing use of systems for recording on plain paper such as electrophotographic and ink jet recording systems, thermal recording becomes often compared with such plain paper recording. For example, thermal recording is required to be close in quality to plain paper recording in terms of stability of recorded portion (image) and stability of unrecorded portion (ground color portion) before and after recording (hereinafter referred to as ground color stability). In particular, as a basic function of thermal recording, the sheet is required to have a good ground color stability to heat and solvents, that is, the sheet is desired to be heat-sensitive only during recording, but non-sensitive in other times.

For the ground color stability of thermal recording sheet, for example, JP-A-4-353490 discloses a thermal recording sheet which undergoes no deterioration in the whiteness of ground color even at high temperatures of about 900°C. This thermal recording sheet contains 4-hydroxydiphenylsulfone and a metal phosphate salt.

Further, rapid increases in the consumption of recording sheets due to increasing construction of various networks, and use of facsimiles and copiers, cause social problems of waste treatment. As a means of solving such problems, a reversible recording material which can be repeatedly recorded and erased is attracting attention.

Reversible recording materials are described in detail in literature such as the Bulletin of the Pulp and Paper Association 47 11 (1993) p1309-1322. For example, JP-A-3-230993 and JP-A-4-366682 disclose recording materials which utilize a reversible change of the recording material between a transparent state and a white turbid state. In addition, recording materials which use reversibility of a thermochromic substance or a reversible color change of a leuco dye have been proposed.

The thermal recording material disclosed in JP-A-OPI 4-353490 shows a Macbeth density of ground color of about 0.11 after heating in a dryer at 950°C for 5 hours, which is a fairly good stability but still insufficient, in term of ground color stability at 120°C.

For the reversible recording materials, one which uses a reversible change of the recording material between a transparent state and a turbid state (in other words, a recording material which utilizes a change in transparency of the recording material) has disadvantages including: (1) image sharpness is poor, (2) clouding (discoloration) speed is low, and (3) when erasing, temperature control is required. Since such a recording material achieves a contrast between transparency and turbidity, there is no problem when it is used in transparent recording sheets such as OHP sheets. However, when it is used in opaque recording sheets such as facsimile paper, it is necessary to provide a coloring layer under the coating layer (the layer capable of reversible change between transparent and turbid states). For this purpose, to provide a contrast in the coating layer, that is, to develop a sharp color of the coloring layer, the coating layer is desirably a thin layer. To make the sheet look white, a thicker layer is preferred. Therefore, the thickness of the coating layer must be strictly controlled.

In reversible recording sheets using a thermochromic material, most thermochromic materials have poor memorizing properties, and thus require a continuous supply of heat to maintain a colour developing condition.

Reversible recording sheets which use leuco dyes as chromogenic sources are disclosed in JP-A-60-193691 and JP-A-257289, but these sheets are erased with water or steam and thus have problems in practical applications. JP-A-2-188293 and JP-A-188294 disclose materials (colour developing/erasing agents) which are simple in structure and have both colour developing and erasing actions to endow a leuco dye with a reversible change in colour by the control of thermal energy.

However, these colour developing/erasing agents involve an erasing process already in the colour developing process, and are thus low in colour developing density. Further, recently, in Japan Hardcopy 93', Yokota et al. reports a recording sheet which uses an amidophenol derivative having a long-chain alkyl group, but this recording sheet requires temperature control during erasing.

An object of the present invention is to provide a thermal recording sheet which has improved ground colour stabil-

ity to heat and solvents. The present invention also aims to provide a thermal recording sheet which has reversible recording capability enabling recording erasing - re-recording.

The above problems are solved by a thermal recording sheet using an urea compound, which is quite different from conventional phenolic colour developers, as a developer.

5 Compounds including urea moieties in their structures are disclosed in, for example, CA-A-1237145. However, these compounds have utility as environmental indicating materials. The document is therefore not relevant to the field of thermal recording sheets.

Thermal recording sheets using urea compound with phenol type developer (Bisphenol A, Bisphenol S, and so on) are disclosed in JP-A-53-140043, 57-87993, 57-82787, and 59-67083. These urea compounds are only limited to the carbon number, and are basically used for depression of melting point of phenol type developer. Thermal recording sheets using urea compound together with a phenol-type developer have a problem of heat resistance, and have no recording reversibility.

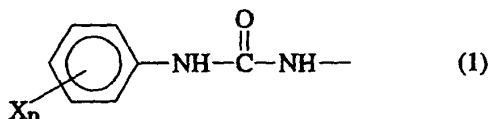
Monourea compounds for use in thermal recording materials are disclosed in JP-A-58-211496, 59-184694, and 61-211085. In these urea compounds only the amino group at one side of the urea is substituted. The compounds thus differ fundamentally from the dimerized and trimerized phenylurea compounds of the present invention.

In general, thermal recording materials are used by dispersing in water, but monourea compounds have a problem since they are slightly soluble in water. Thermal recording materials using these monourea compounds alone have a problem of heat resistance, and have no reversible recording.

Dimerized urea-type compounds are disclosed in JP-A-5-1317152 and 5-147357. These dimerized urea-type compounds are characterised by a structure having a sulfonyl group adjacent to the urea structure ($\text{Ar-SO}_2\text{-NH-C(=O)-NH-}$). However, the thermal recording sheets using these dimerized urea also have a problem in heat resistance, and have no reversible recording.

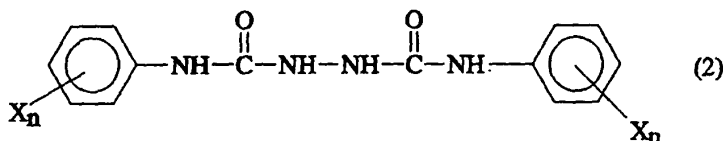
EP-A-0615859 and EP-A-0620122 describe thermosensitive recording materials in which the colour developing agent comprises an aromatic urea or thiourea derivative comprising the divalent structural unit $\text{-SO}_2\text{NHC(=X)NH-}$ wherein X is oxygen or sulphur. These two documents belong to the state of the art according to Article 54(3)EPC for the commonly designate states DE and GB.

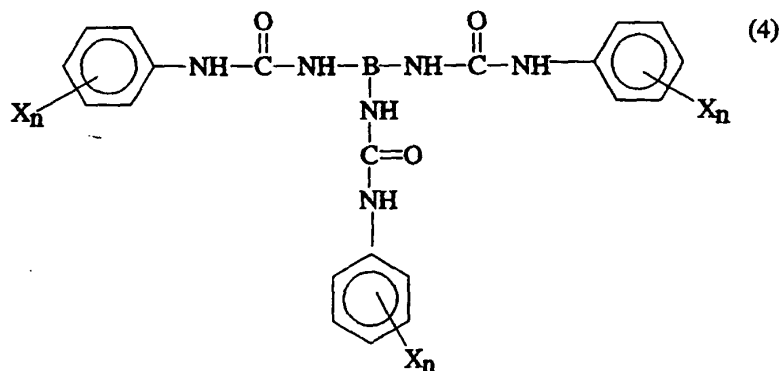
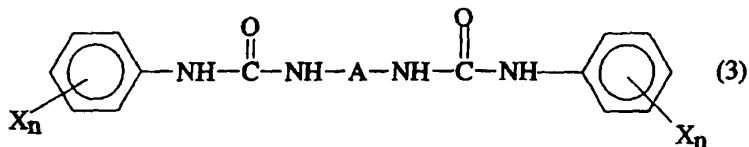
The present invention provides a thermal recording sheet which comprises a thermal recording layer comprising a colourless or pale coloured dye precursor and a colour developer, which colour developer is a urea compound which comprises at least two groups of formula (1)



wherein X is $\text{C}_1\text{-C}_{12}$ alkyl, halogenated $\text{C}_1\text{-C}_6$ alkyl, $\text{C}_7\text{-C}_{14}$ aralkyl, $\text{C}_1\text{-C}_{12}$ alkoxy, $\text{C}_6\text{-C}_{12}$ aryloxy, $\text{C}_1\text{-C}_{12}$ alkoxycarbonyl, $\text{C}_1\text{-C}_{12}$ acyl, $\text{C}_1\text{-C}_{12}$ dialkylamino, $\text{C}_7\text{-C}_{12}$ arylalkylamino, $\text{C}_6\text{-C}_{12}$ arylamino, $\text{C}_1\text{-C}_{12}$ acylamino, nitro, cyano, a halogen or hydrogen, and n is an integer of 1 to 3. The colour developer reacts with the dye precursor upon heating to develop a colour.

In a preferred aspect of the invention, the thermal recording sheet comprises as a colour developer a urea compound of Formula (2), (3), or (4)

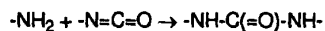




30 wherein X is as defined above for formula (1), A is a divalent group comprising 30 or less carbon atoms, and B is a trivalent group comprising 30 or less carbon atoms and n is an integer from 1 to 3.

The group of formula (1) is hereinafter referred to as the phenylurea structure. A urea compound of Formula (2), (3), or (4) is a compound having two or three phenylurea structures of Formula (1). A urea compound of Formula (2) is a compound in which two phenylurea structures are directly bonded, and a urea compound of Formula (3) or (4) is a compound in which two or three phenylurea structures are bonded through a joint group. The joint group is a divalent or trivalent group comprising 30 or less carbon atoms.

35 An urea compound of Formula (2) or (3) (hereinafter referred to as a "dimerized urea compound") and a urea compound of Formula (4) (hereinafter referred to as a "trimerized urea compound") can be individually synthesized, for example, by a reaction of amines with an isocyanate compound.



First, the dimerized urea compound of Formula (3) can be synthesized (a) by reacting diamines with a monoisocyanate compound, or (b) by reacting a diisocyanate compound with monoamines.

45 (a)



50 (b)



When diamines are reacted with a monoisocyanate compound, the diamines include carbonyldiamide, ethylenediamine, 1,2-propanediamine, 1,3-propanediamine, 1,4-diaminobutane, 1,3-diaminopentane, 1,5-diaminopentane, 1,6-diaminohexane, 1,7-diaminoheptane, 1,8-diaminooctane, 1,9-diaminononane, 1,10-diaminodecane, 1,11-diaminoundecane, 1,12-diaminododecane, 1,2-diamino-2-methylpropane, 1,3-diamino-2,2-dimethylpropane, 1,5-diamino-2-methylpentane, 1,5-diamino-2,2-dimethylpentane, trimethylhexamethylenediamine, diethylenetriamine, triethylenetetra-

tramine, tetraethylenepentamine, 3,3'-diaminodipropylamine, N,N-bis(3-aminopropyl)methylamine, bis(3-aminopropyl)ether, 1,8-diamino-3,6-dioxaoctane, ethyleneglycolbis(3-aminopropyl)ether, 1,2-diaminocyclohexane, 1,3-diaminocyclohexane, 1,4-diaminocyclohexane, 1,3-di(aminomethyl)cyclohexane, 1,4-di(aminomethyl)cyclohexane, 4,4'-diaminodiphenylsulfone, 3,3'-diaminodiphenylsulfone, 4,4'-diaminodiphenylsulfide, 4,4'-diaminodiphenylether, 3,4'-diaminodiphenylether, 4,4'-diaminodiphenylether, 3,4'-diaminodiphenylether, 4,4'-diaminodicyclohexylmethane, 4,4'-diaminodiphenylmethane, 3,3'-diaminodiphenylmethane, 4,4'-diamino-3,3'-dichlorodiphenylmethane, 4,4'-diamino-1,2-diphenylethane, 4,4'-diamino-2,2'-dimethylbiphenyl, 4,4'-diaminodiphenylamine, 2,2'-diaminodiphenyldisulfide, 4,4'-diaminodiphenyldisulfide, 1,3-bis(m-aminophenoxy)benzene, 1,4-bis(m-aminophenoxy)benzene, 2,2-bis(4-aminophenyl)propane, 2,2-bis(4-(4-aminophenoxy)phenyl)hexafluoropropane, a,a'-bis(4-aminophenyl)-1,4-diisopropylbenzene, 3,3'-diaminobenzophenone, 4,4'-diaminobenzophenone, 2,4-diaminoazobenzene, 4,4'-diaminostilbene, 4,4'-diaminostilbene-2,2'-disulfonic acid, o-phenylenediamine, m-phenylenediamine, p-phenylenediamine, 4-chloro-o-phenylenediamine, 5-chloro-m-phenylenediamine, 2-chloro-p-phenylenediamine, 2-nitro-1,4-phenylenediamine, 4-nitro-1,2-phenylenediamine, 4-nitro-1,3-phenylenediamine, 2,4-diaminoanisole, p,p'-methylenedianiline, m-xylylenediamine, p-xylylenediamine, 2,4-diaminotoluene, 2,6-diaminotoluene, 3,4-diaminobenzoic acid, 3,5-diaminobenzoic acid, 1,5-diaminonaphthalene, 1,8-diaminonaphthalene, 2,3-dianonaphthalene, 1,2-diaminoanthraquinone, 1,4-diaminoanthraquinone, 1,5-diaminoanthraquinone, 2,6-diaminoanthraquinone, 1,4-diamino-2,3-dicyano-9,10-anthraquinone, 2,3-diaminopyridine, 2,6-diaminopyridine, 3,4-diaminopyridine, 2,4-diamino-6-chloropyrimidine, 2,4-diamino-6-phenyl-1,3,5-triazine, 9,9-bis(4-aminophenyl)fluorene, piperazine, N-aminoethylpiperazine, 1,4-bis(3-aminopropyl)piperazine, and isophoronediamine; the monoisocyanate compound includes aromatic isocyanate compounds such as phenylisocyanate, o-tolylisocyanate, m-tolylisocyanate, p-tolylisocyanate, 3,4-dimethylphenylisocyanate, 2,6-dimethylphenylisocyanate, 3,4,5-trimethylphenylisocyanate, o-trifluoromethylphenylisocyanate, m-trifluoromethylphenylisocyanate, p-trifluoromethylphenylisocyanate, 2-methoxyphenylisocyanate, 3-methoxyphenylisocyanate, 4-methoxyphenylisocyanate, p-bromophenylisocyanate, o-fluorophenylisocyanate, m-fluorophenylisocyanate, p-fluorophenylisocyanate, o-chlorophenylisocyanate, 2,5-dichlorophenylisocyanate, 3,4-dichlorophenylisocyanate, 2,6-dichlorophenylisocyanate, o-nitrophenylisocyanate, m-nitrophenylisocyanate, p-nitrophenylisocyanate, p-dimethylaminophenylisocyanate, and p-diethylaminophenylisocyanate, and any combinations of both compounds can be used.

On the other hand, when a diisocyanate compound is reacted with monoamines, the diisocyanate compound includes 2,4-toluenediisocyanate (2,4-TDI), 2,6-toluenediisocyanate (2,6-TDI), 4,4'-diphenylmethanediisocyanate (MDI), 1,5-naphthalenediisocyanate (NDI), toluenediisocyanate (TODI), hexamethylenediisocyanate (HDI), isophoronediiisocyanate (IPDI), p-phenylenediisocyanate, m-phenylenediisocyanate, trans-cyclohexane-1,4-diisocyanate, xylylenediisocyanate (XDI), 4,4'-dicyclohexylmethanediisocyanate (hydrogenated MDI), lysinediisocyanate (LDI), m-tetramethylxylenediisocyanate (m-TMXDI), and p-tetramethylxylenediisocyanate (p-TMXDI); the monoamines include aromatic amines such as aniline, o-toluidine, m-toluidine, p-toluidine, o-ethylaniline, m-ethylaniline, p-ethylaniline, p-n-propylaniline, p-iso-propylaniline, p-n-butylaniline, 2-sec-butylaniline, 4-sec-butylaniline, 4-tert-butylaniline, p-n-amylianiline, p-n-hexylaniline, p-n-heptylaniline, p-n-octylaniline, p-n-nonylaniline, p-n-decylaniline, p-undecylaniline, p-dodecylaniline, 2,3-xylylidine, 2,4-xylylidine, 2,5-xylylidine, 3,4-xylylidine, mesitylamine, 2,6-diethylaniline, o-aminobenzotrifluoride, m-aminobenzotrifluoride, p-aminobenzotrifluoride, o-methoxyaniline, m-methoxyaniline, p-methoxyaniline, o-ethoxyaniline, m-ethoxyaniline, p-ethoxyaniline, 4-tert-butoxyaniline, 2,4-dimethoxyaniline, 2,5-dimethoxyaniline, 3,4-dimethoxyaniline, 3,5-dimethoxyaniline, o-chloroaniline, m-chloroaniline, p-chloroaniline, o-bromoaniline, m-bromoaniline, p-bromoaniline, o-fluoroaniline, m-fluoroaniline, p-fluoroaniline, 2,3-dichloroaniline, 2,4-dichloroaniline, 2,5-dichloroaniline, 2,6-dichloroaniline, 3,4-dichloroaniline, 3,5-dichloroaniline, 2,4-dibromoaniline, 2,5-dibromoaniline, 2,6-dibromoaniline, 2,4-difluoroaniline, 2,5-difluoroaniline, 2,6-difluoroaniline, 3,4-difluoroaniline, 2,3,4-trichloroaniline, 2,4,5-trichloroaniline, 2,4,6-trichloroaniline, o-nitroaniline, m-nitroaniline, p-nitroaniline, o-cyanoaniline, m-cyanoaniline, p-cyanoaniline, methyl o-aminobenzoate, methyl m-aminobenzoate, methyl p-aminobenzoate, ethyl o-aminobenzoate, ethyl m-aminobenzoate, ethyl p-aminobenzoate, n-propyl p-aminobenzoate, iso-propyl p-aminobenzoate, n-butyl p-aminobenzoate, iso-butyl p-aminobenzoate, phenyl o-aminobenzoate, phenyl m-aminobenzoate, phenyl p-aminobenzoate, benzyl o-aminobenzoate, benzyl m-aminobenzoate, benzyl p-aminobenzoate, p-nitrophenyl o-aminobenzoate, 2-diethylaminoethyl p-aminobenzoate, 2'-aminoacetophenone, 3'-aminoacetophenone, 4'-aminoacetophenone, 2-aminobenzophenone, 3-aminobenzophenone, 4-aminobenzophenone, m-aminobenzoylmethylamide, 2-aminodiphenylether, 3-aminodiphenylether, 4-aminodiphenylether, 2-amino-4'-chlorodiphenylether, 4-amino-4'-chlorodiphenylether, 4-amino-4'-aminodiphenylmethane, 4-amino-4'-chlorodiphenylmethane, 2-amino-5-chlorotoluene, 4-amino-2-chlorotoluene, 2-amino-5-boromobenzotrifluoride, 2-amino-5-chlorobenzotrifluoride, 3-amino-4-chlorobenzotrifluoride, 5-amino-2-chlorobenzotrifluoride, and any combinations of both compounds can be used.

The dimerized urea compound of Formula (2) can be synthesized using hydrazine as the diamine and the above compounds as the monoisocyanate compound. On the other hand, the trimerized urea compound of Formula (4) can be synthesized by a method of reacting triamines with a monoisocyanate compound or a method of reacting a triisocyanate compound with monoamines.

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First, when triamines are reacted with a monoisocyanate compound, the triamines can be 1,2,4-triaminobenzene, tris(2-aminoethyl)amine, melamine, 2,4,6-triaminopyrimidine, and triamtherene, the monoisocyanate compound can be the above described compounds, and any combinations of both compounds can be used. When a triisocyanate compound is reacted with monoamines, the triisocyanate compound can be triphenylmethanetriisocyanate, tris(isocyanatephenyl)thiophosphate, lysine ester triisocyanate, 1,6,11-undecanetriisocyanate, 1,8-diisocyanate-4-
5 isocyanatemethyloctane, 1,3,6-hexamethylenetriisocyanate, and bicycloheptanetriisocyanate; the monoamines can be the above described compounds, and any combinations of both compounds can be used.

Practical examples of the urea compound of Formula (2), (3), or (4) are shown below:

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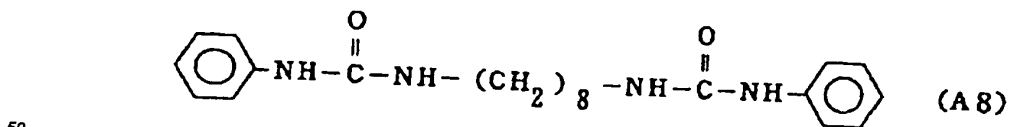
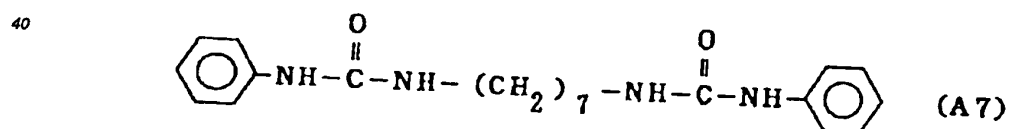
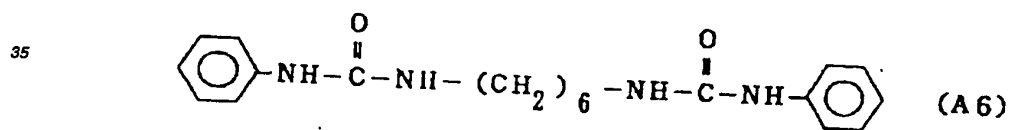
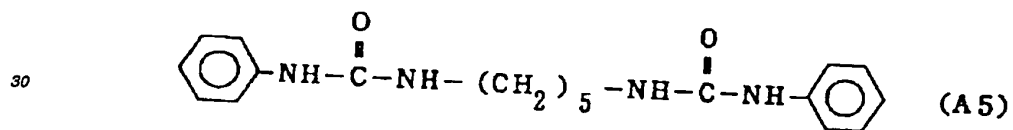
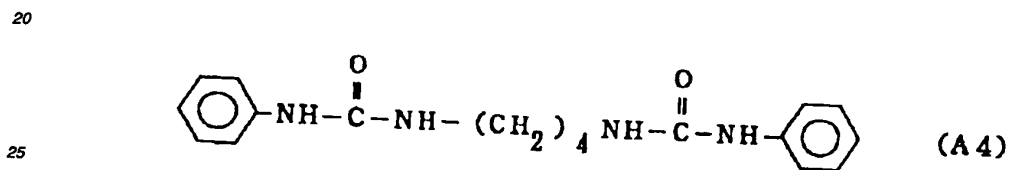
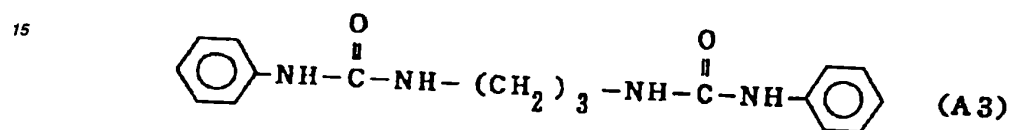
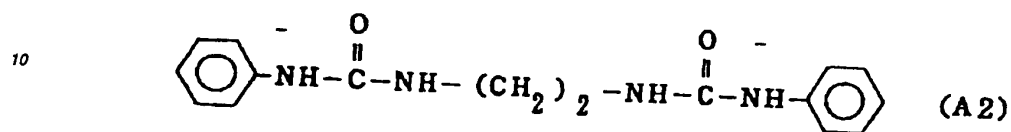
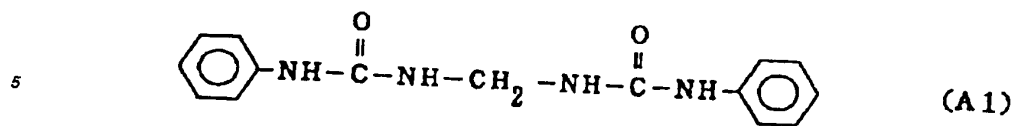
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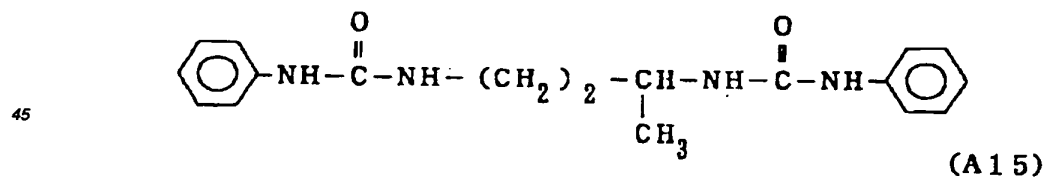
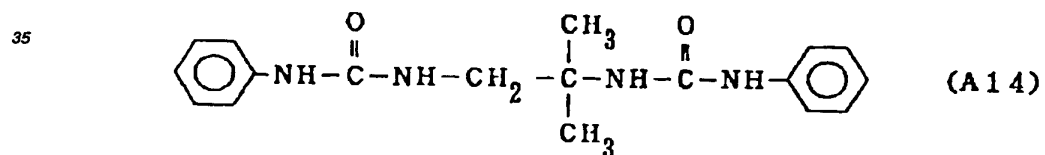
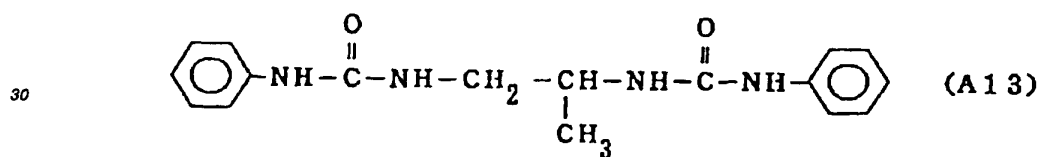
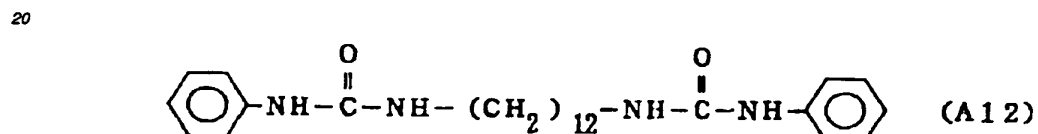
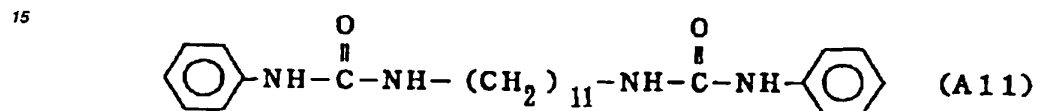
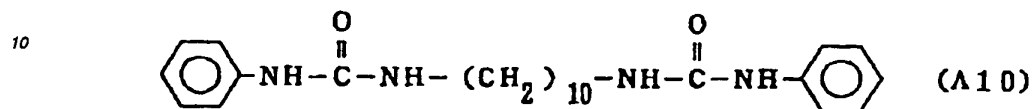
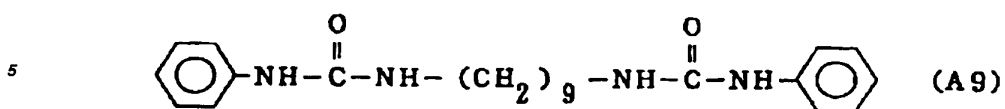
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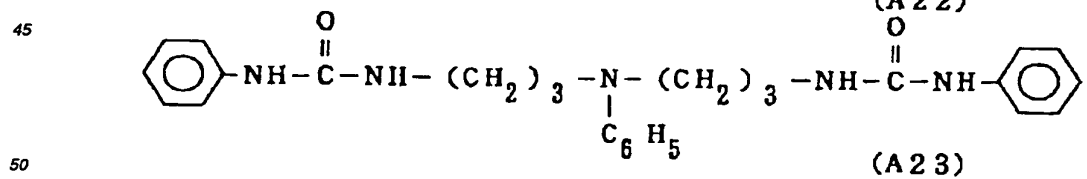
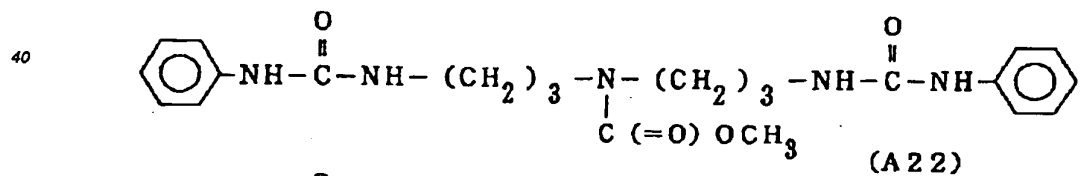
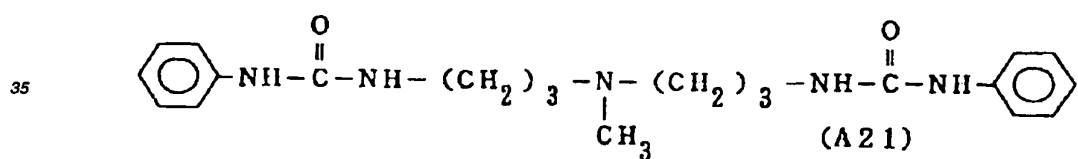
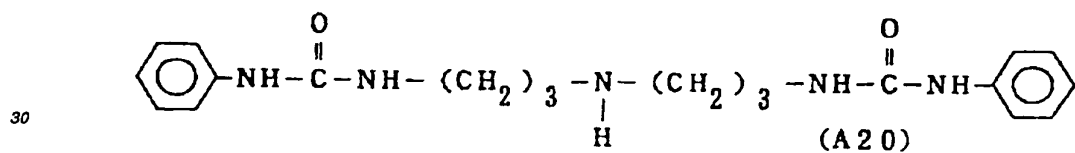
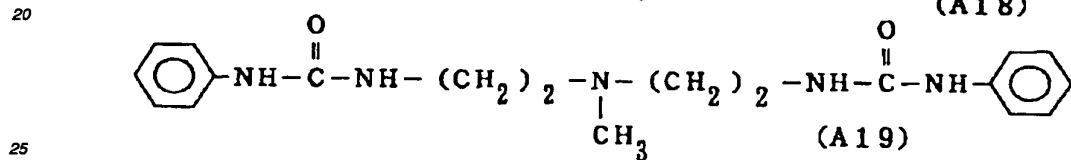
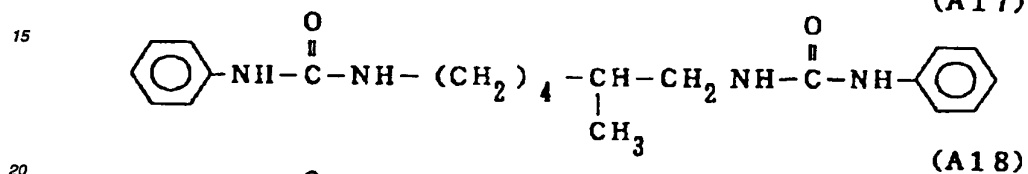
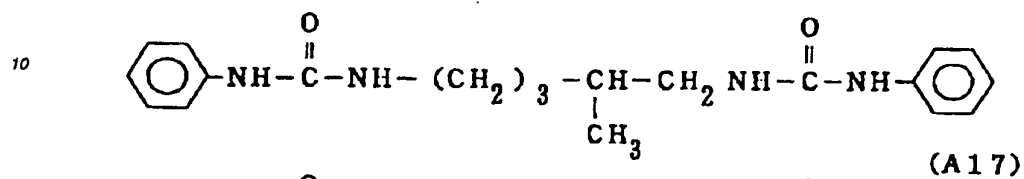
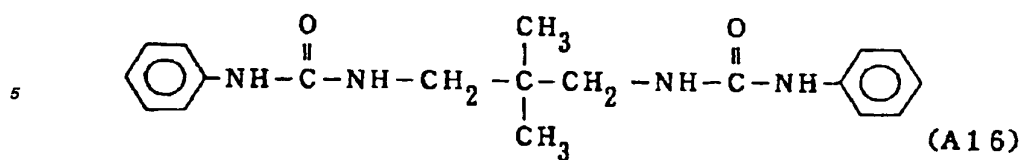
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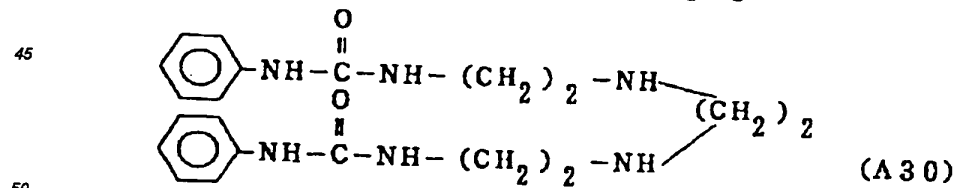
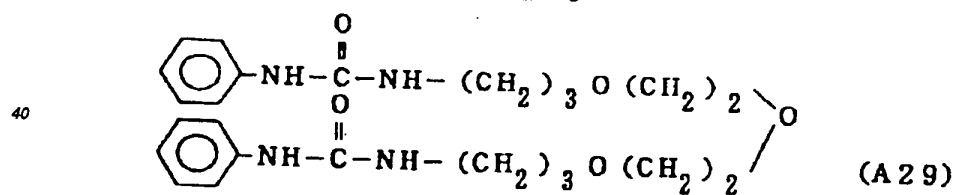
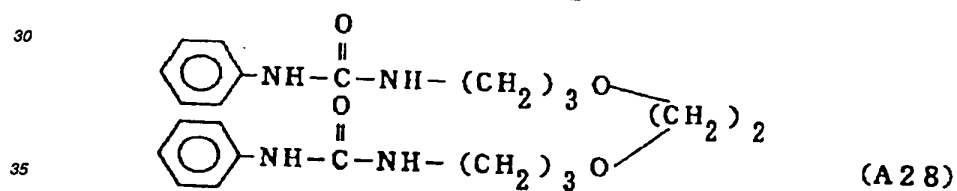
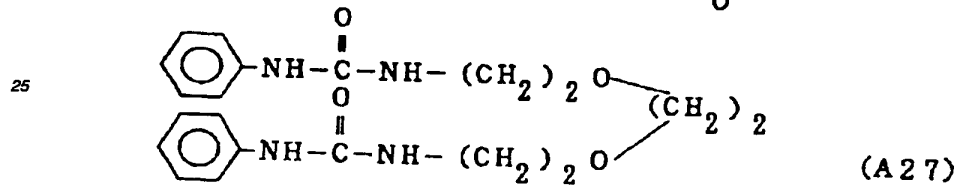
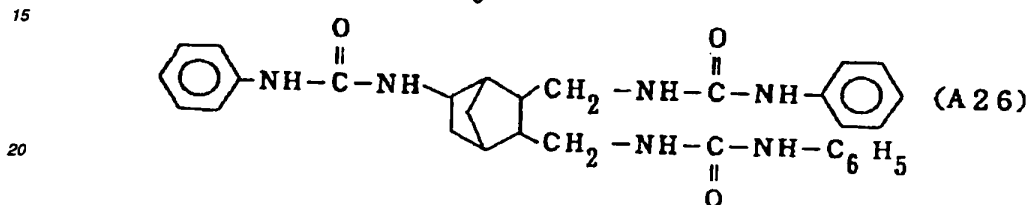
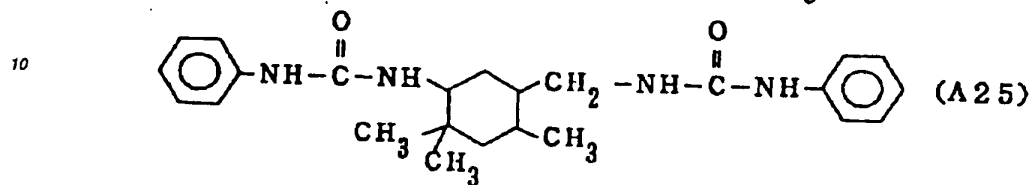
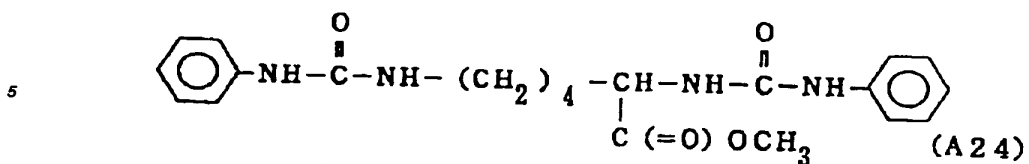


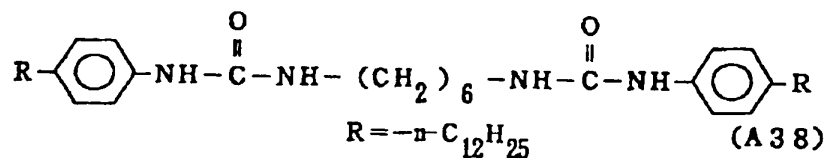
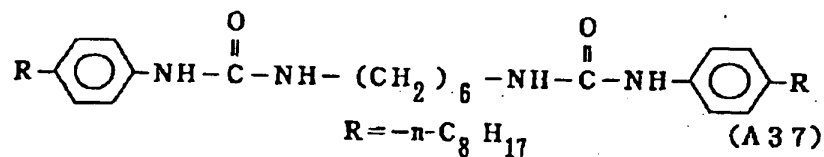
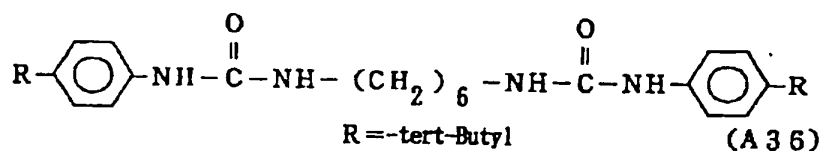
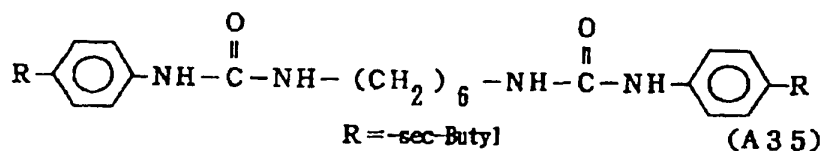
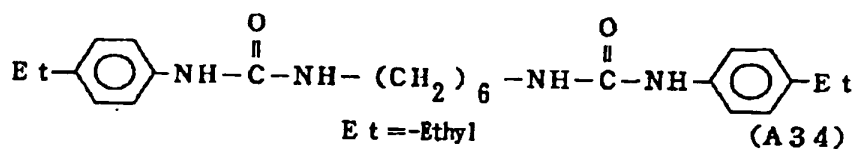
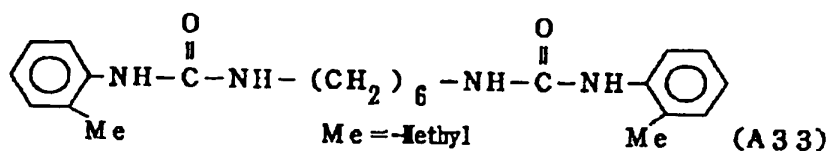
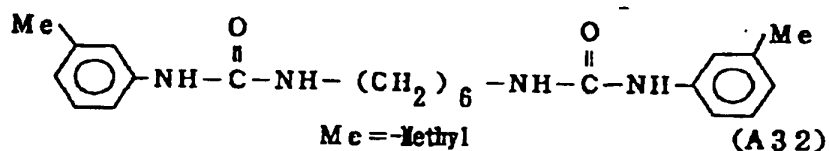
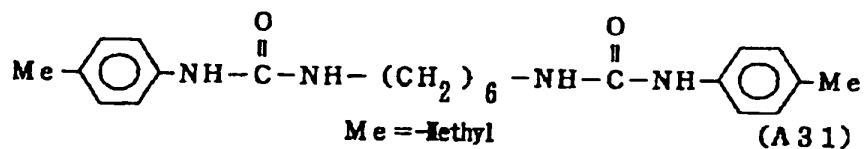


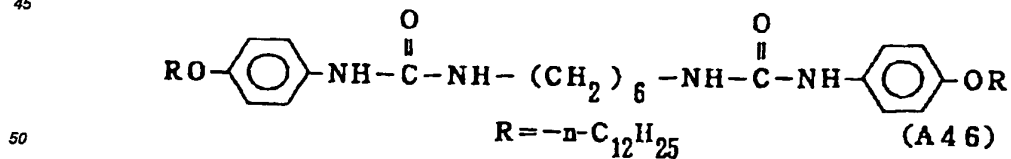
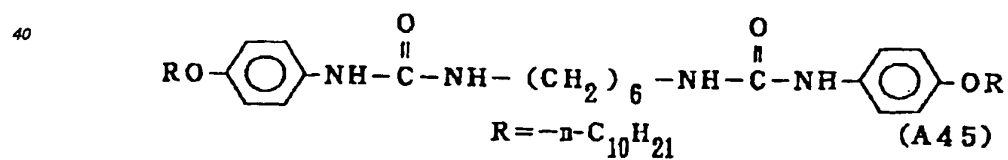
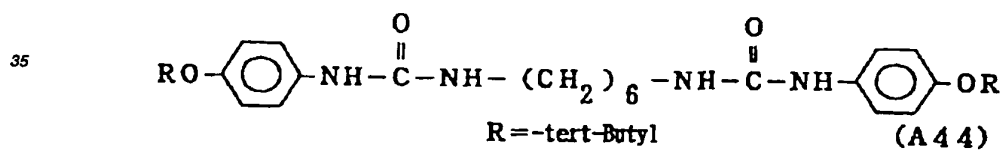
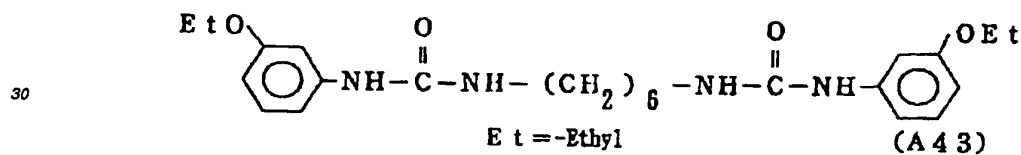
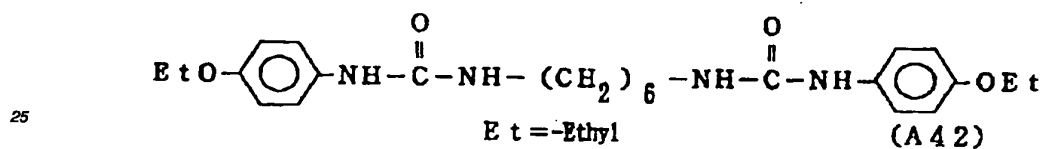
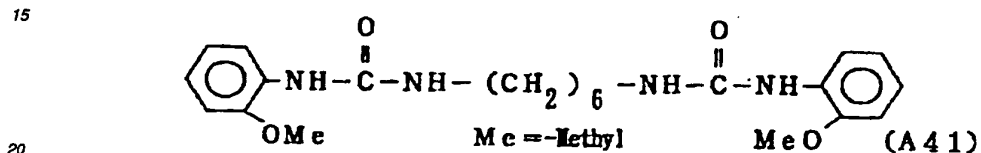
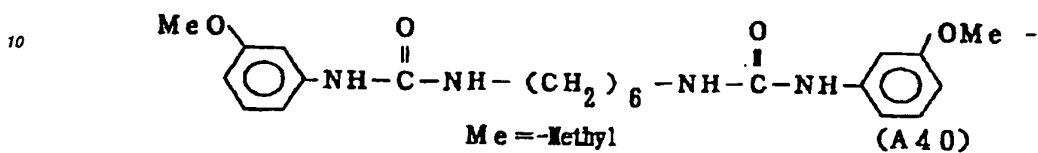
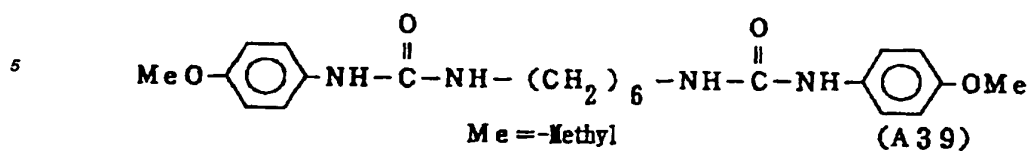
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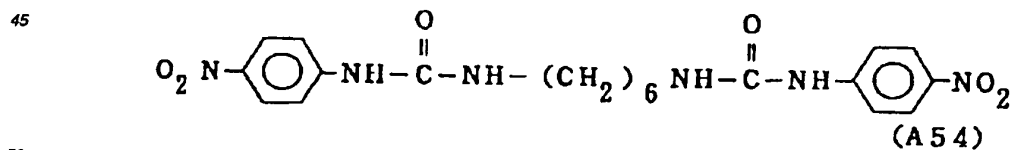
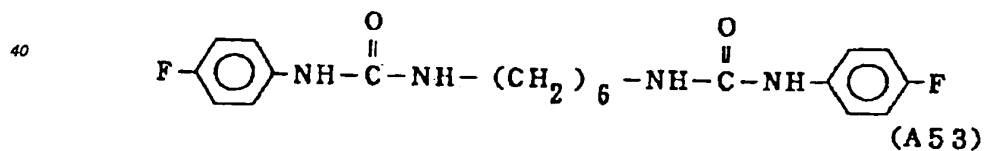
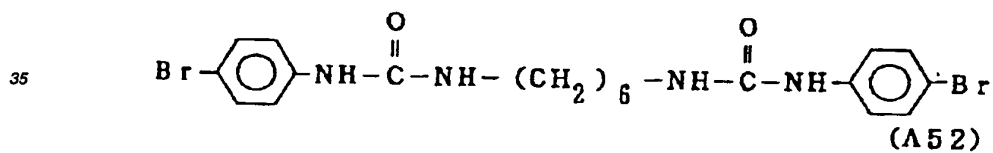
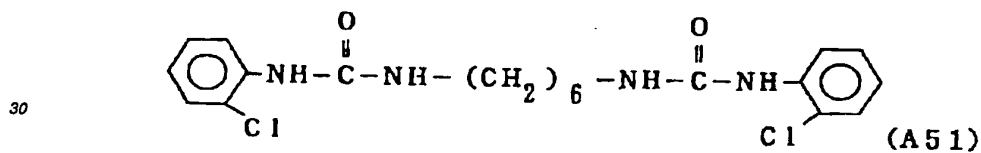
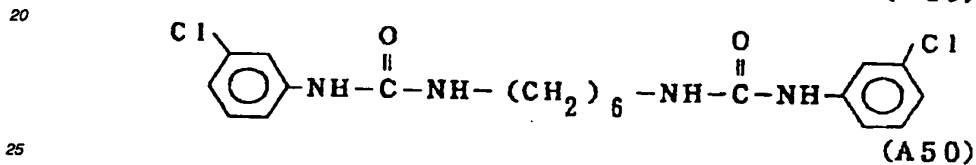
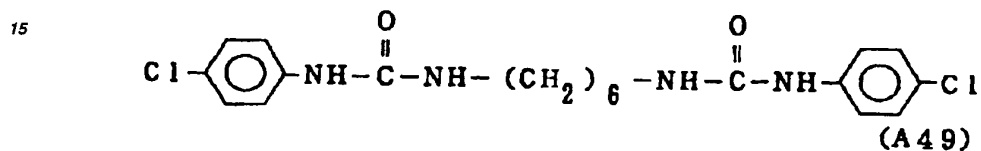
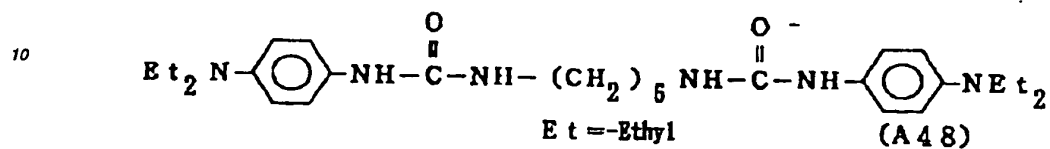
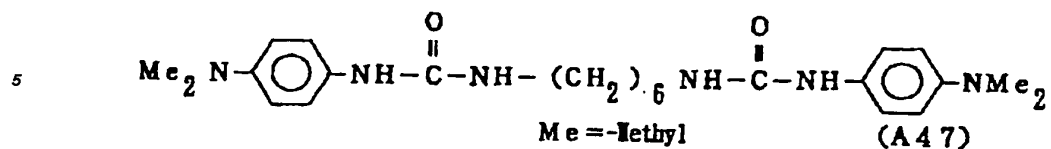
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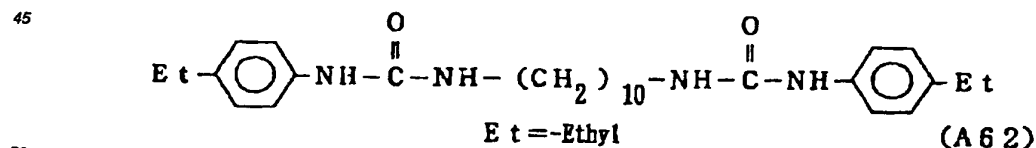
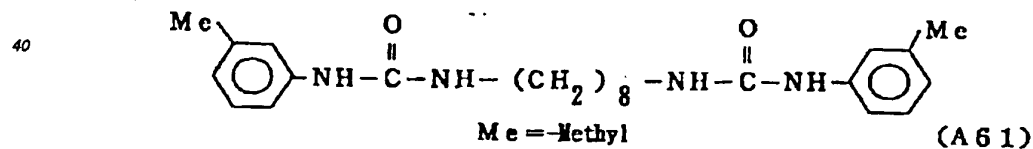
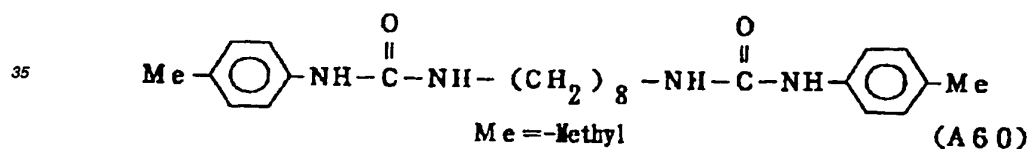
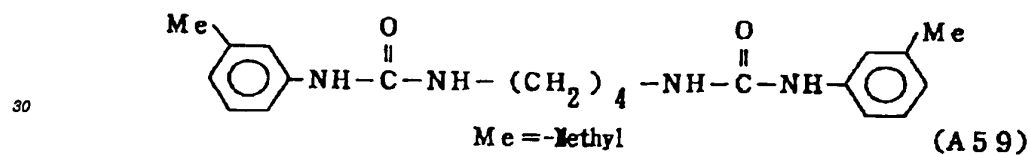
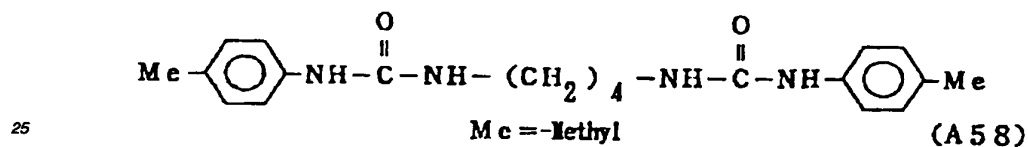
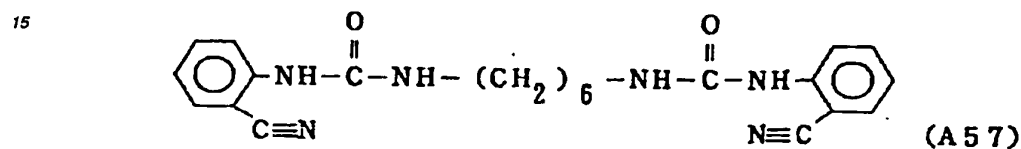
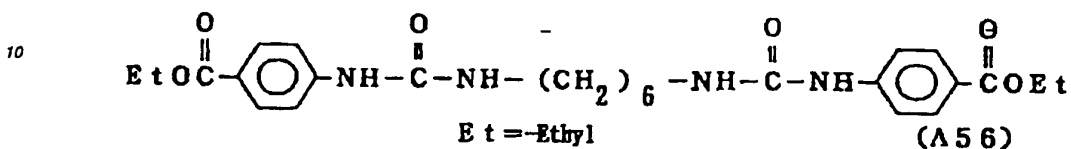
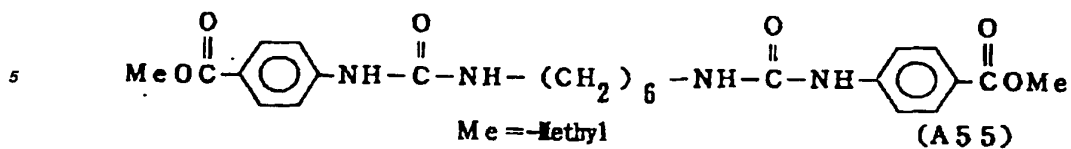


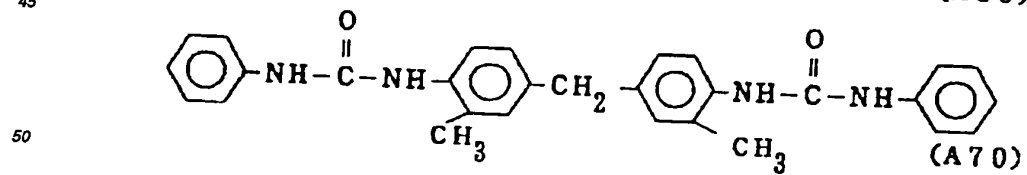
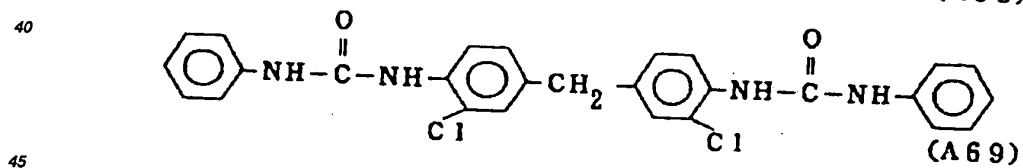
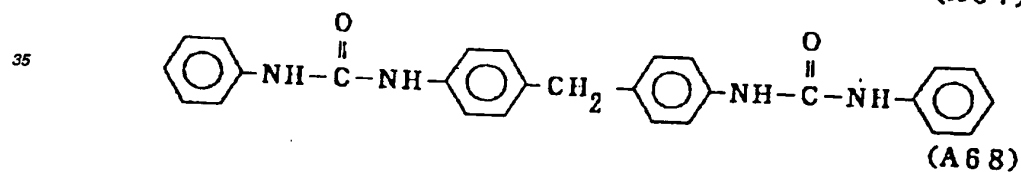
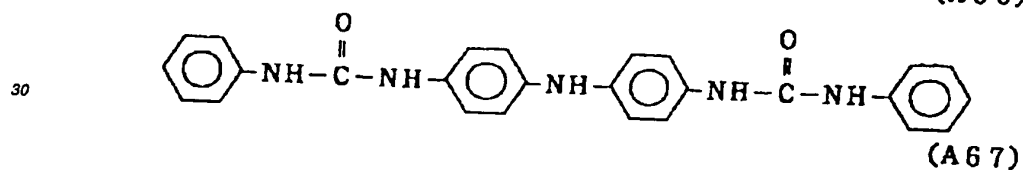
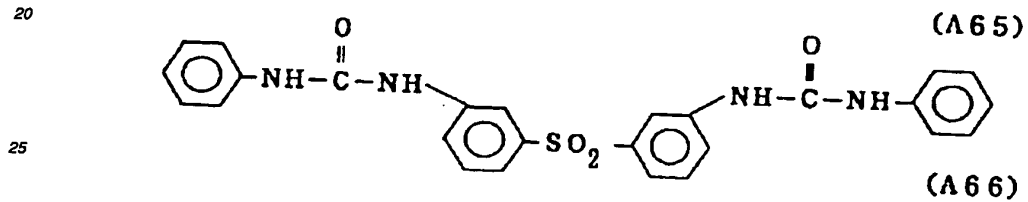
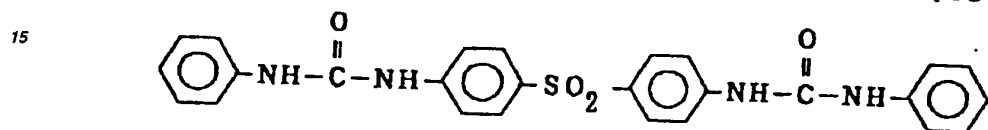
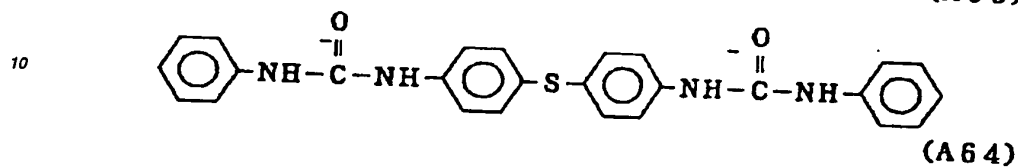
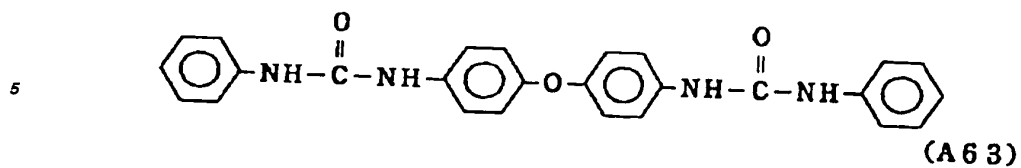


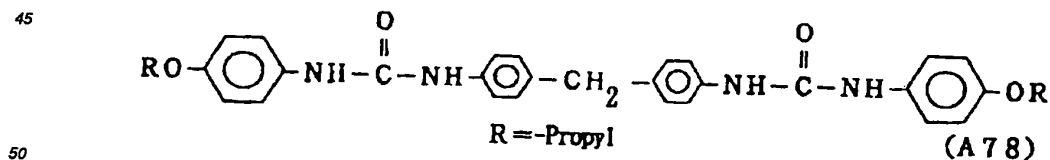
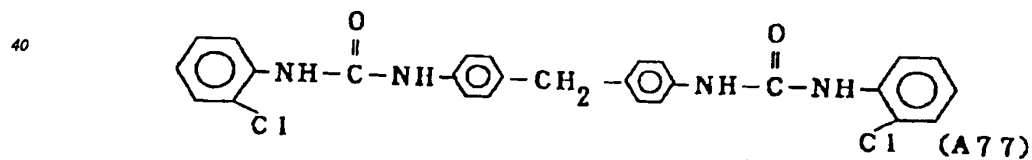
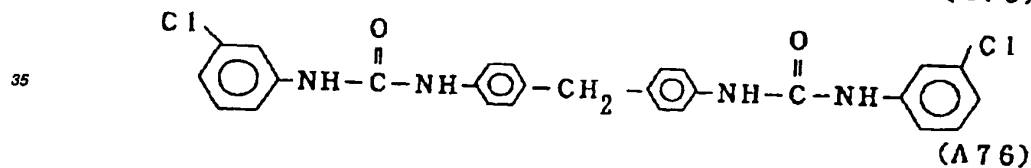
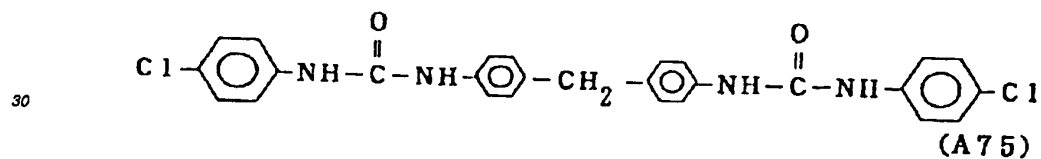
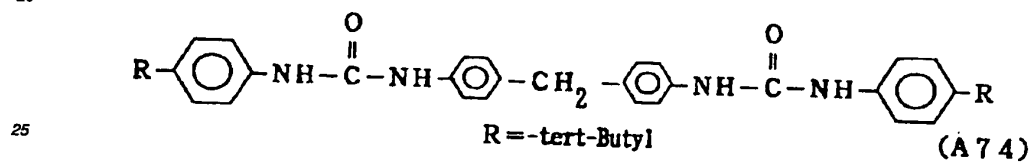
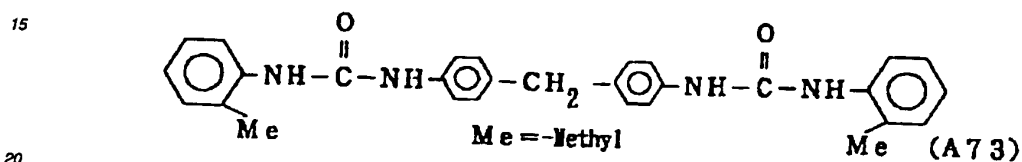
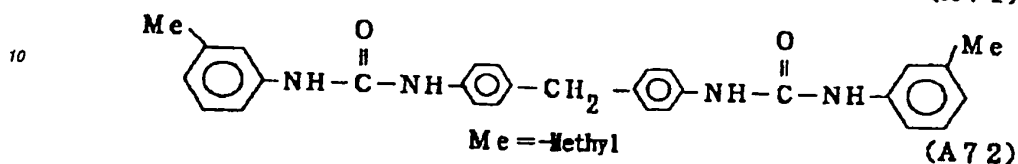
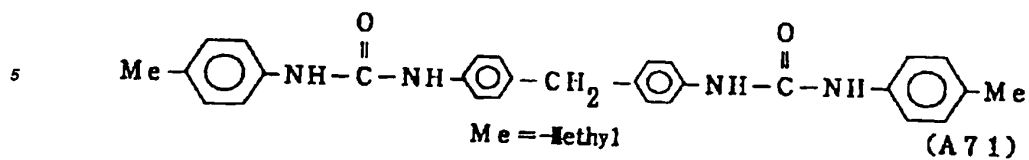






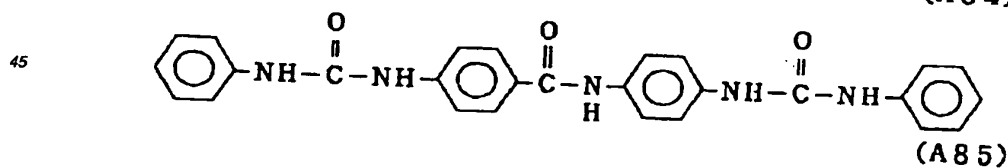
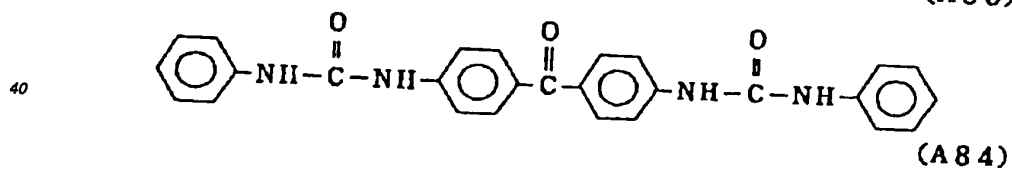
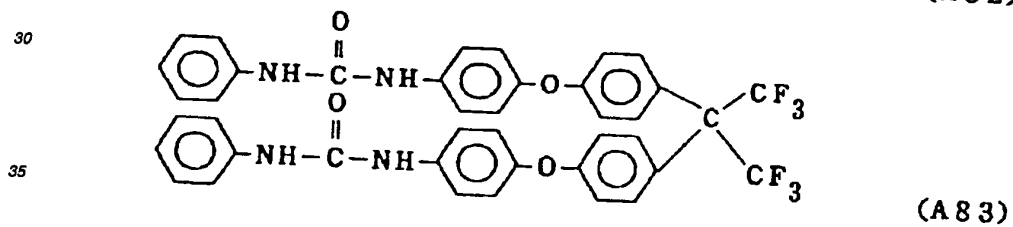
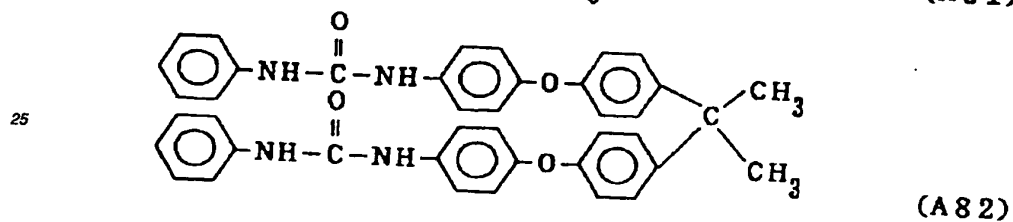
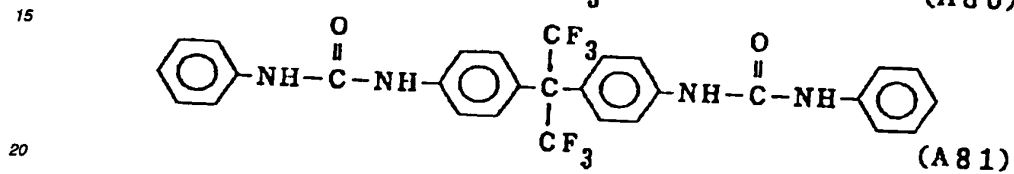
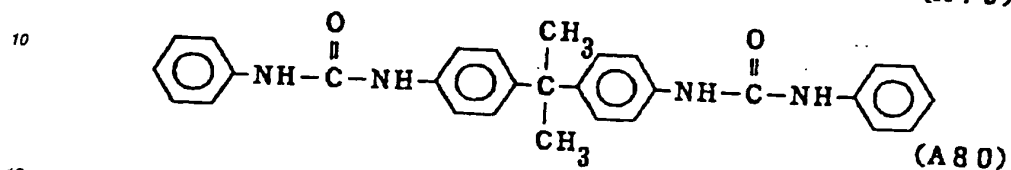
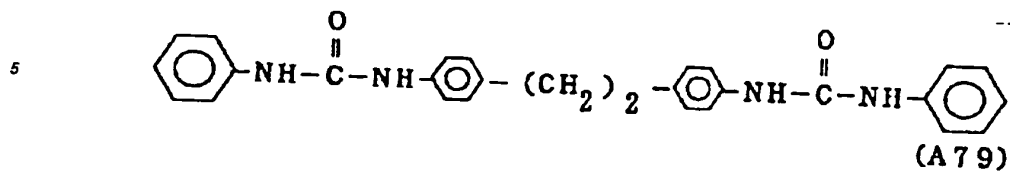






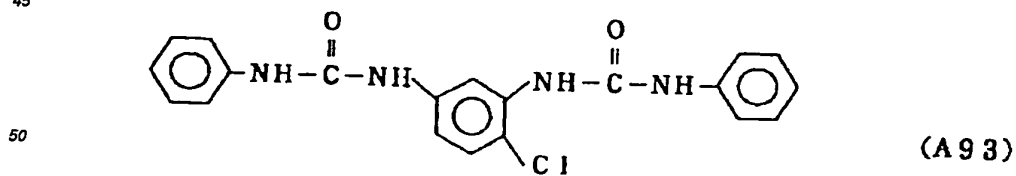
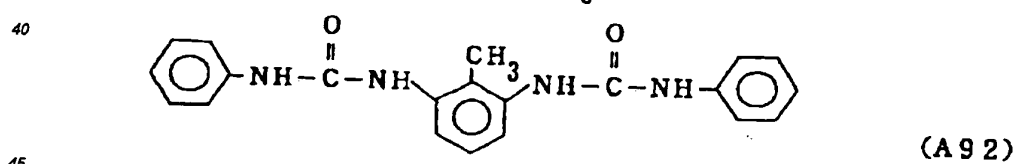
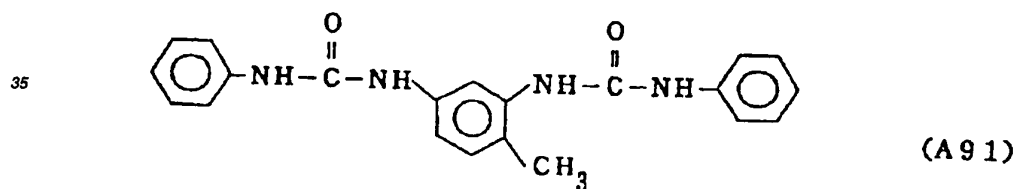
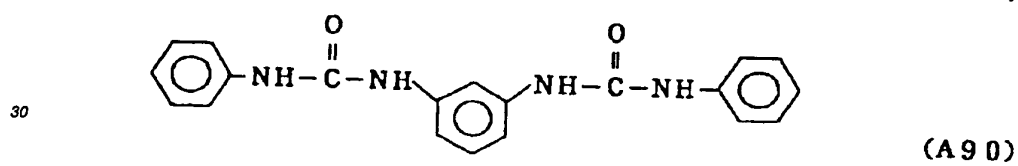
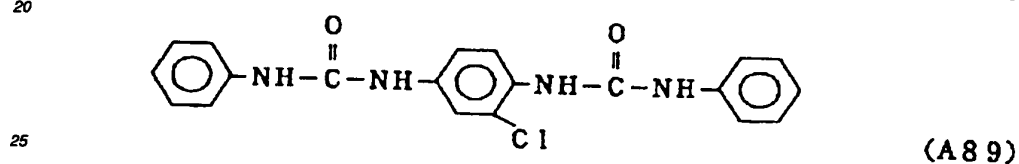
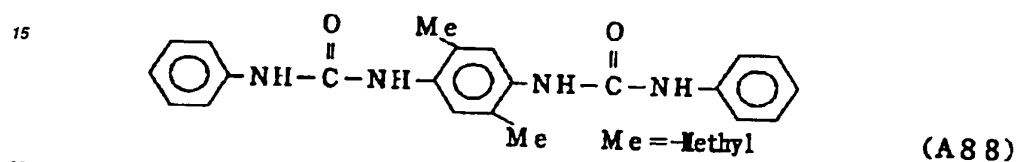
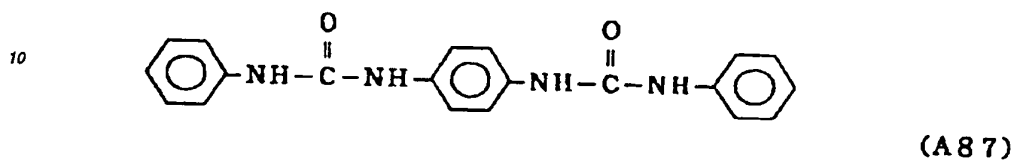
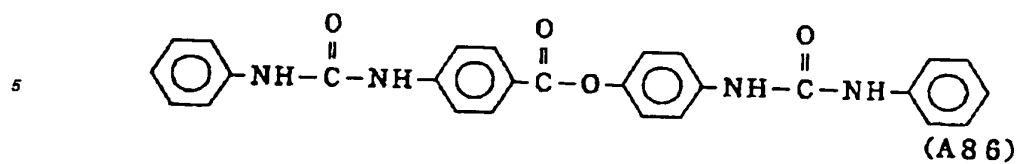
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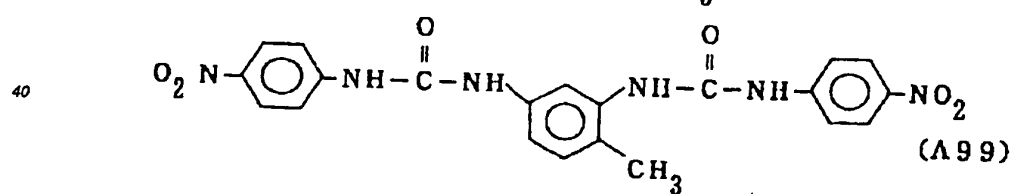
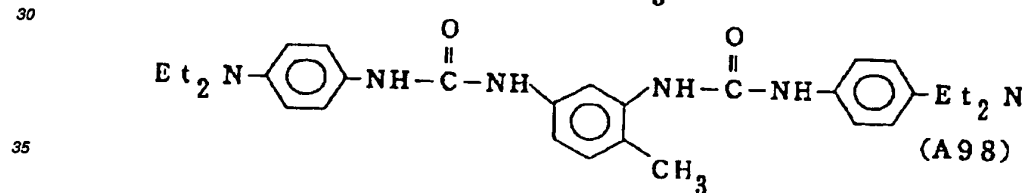
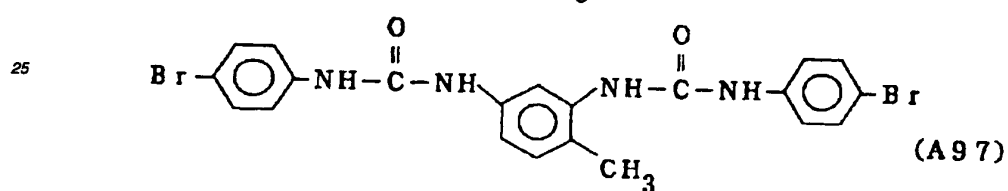
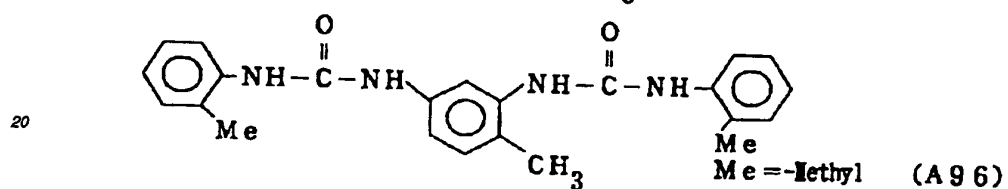
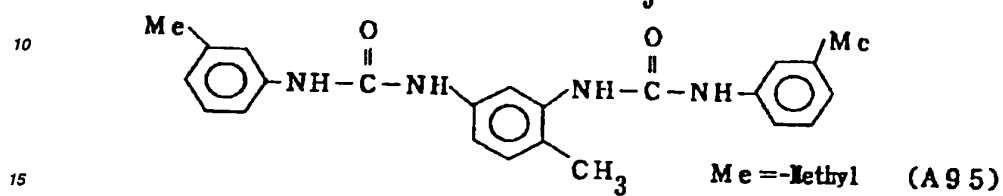
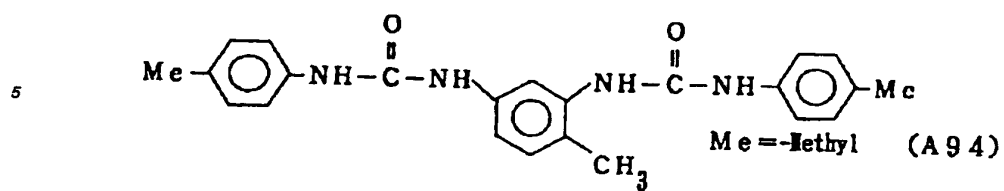
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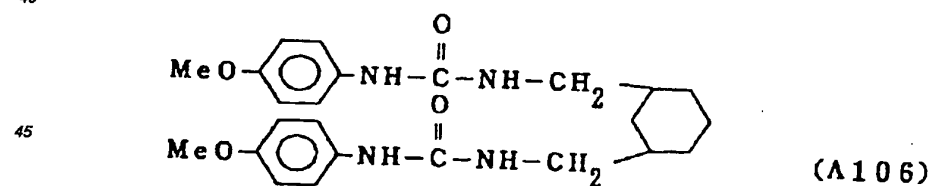
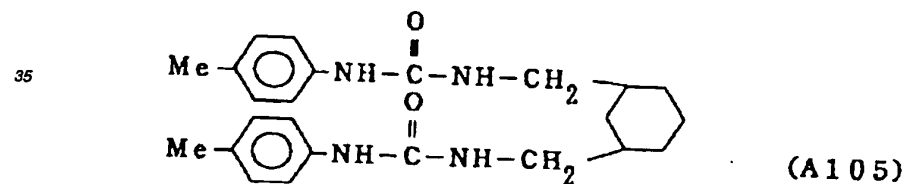
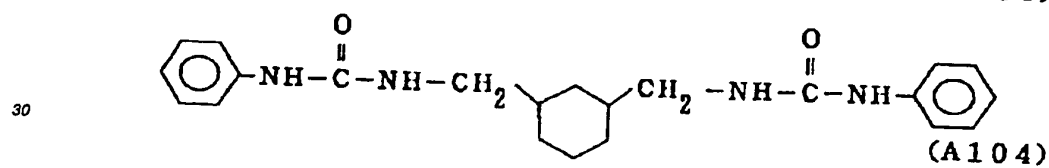
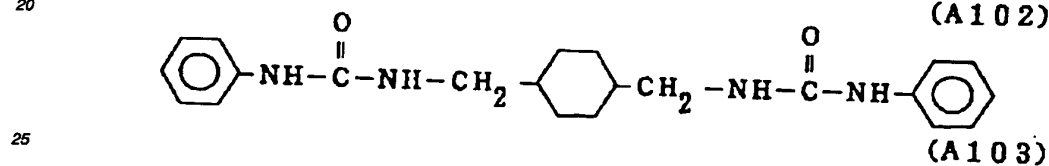
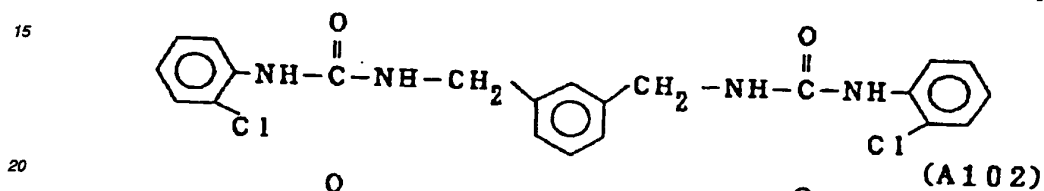
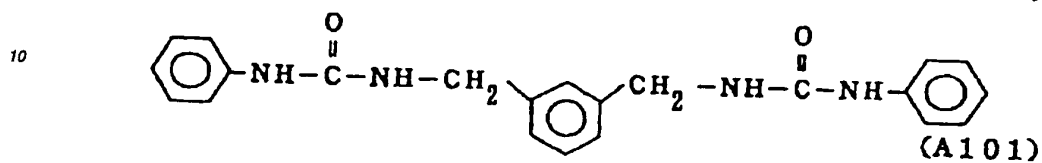
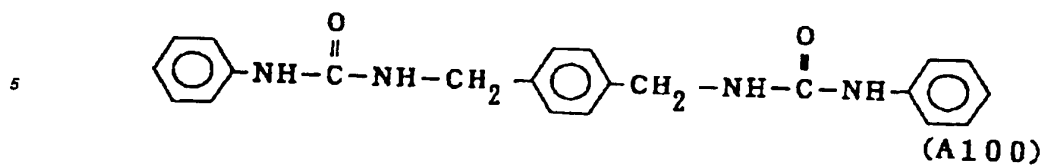




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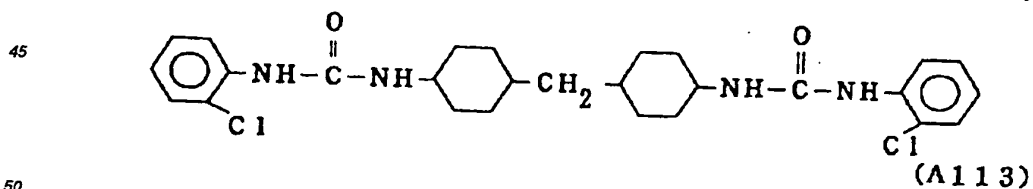
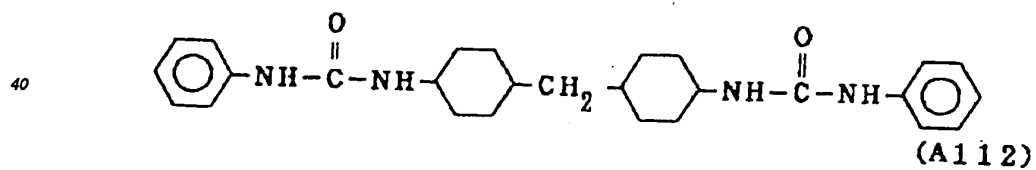
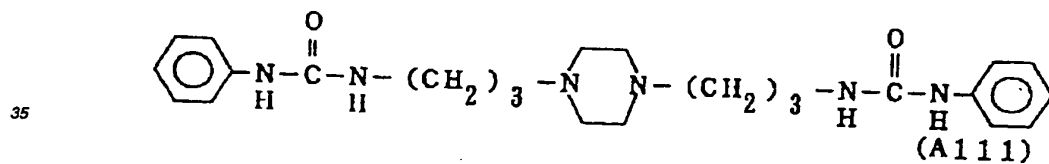
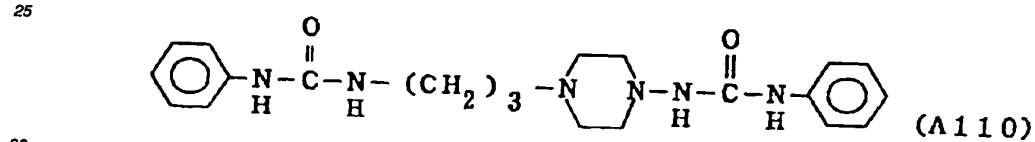
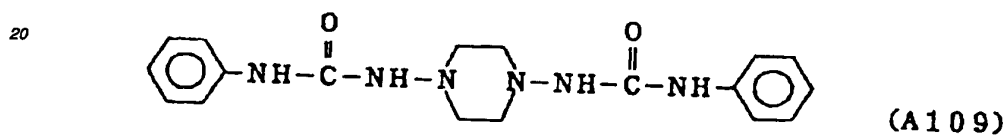
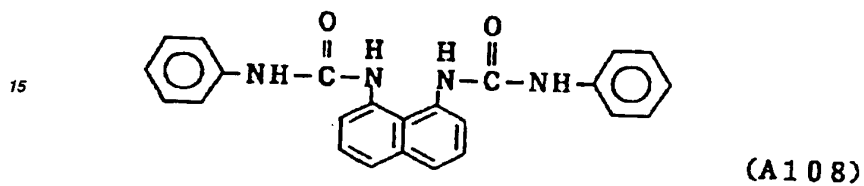
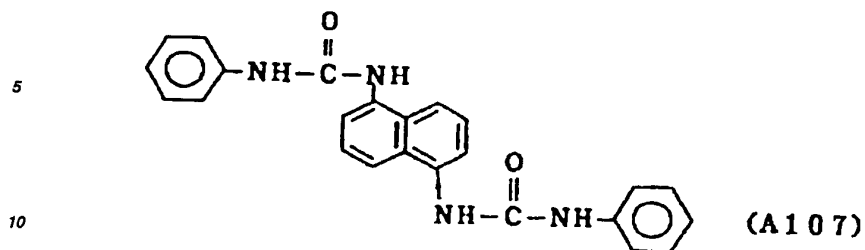
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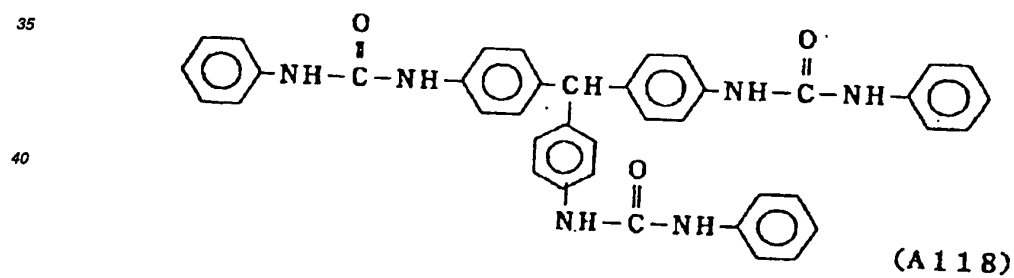
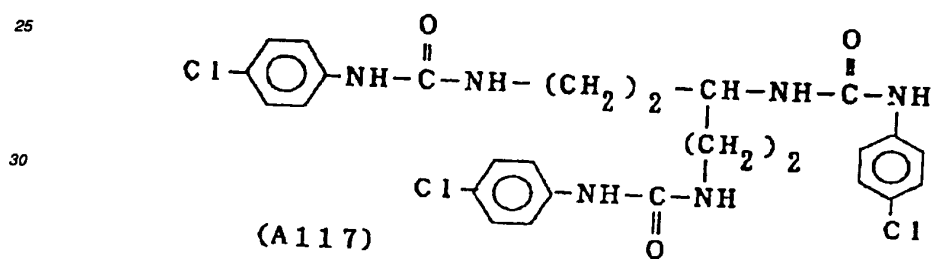
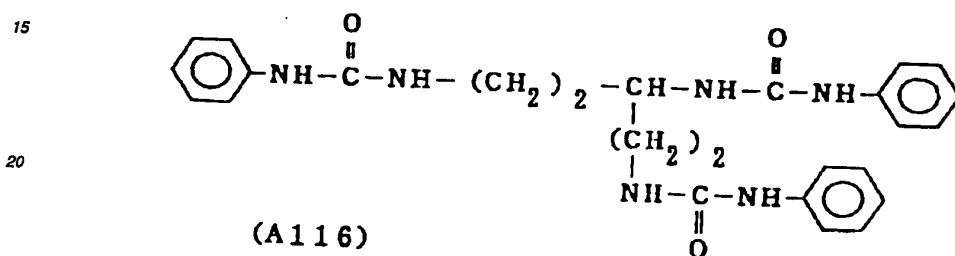
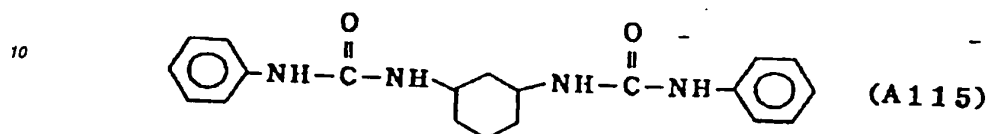
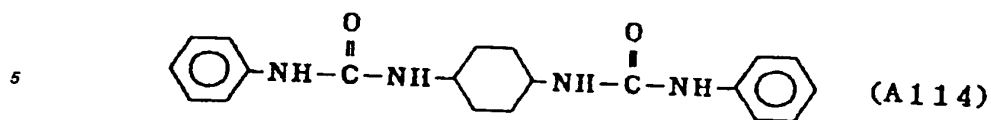
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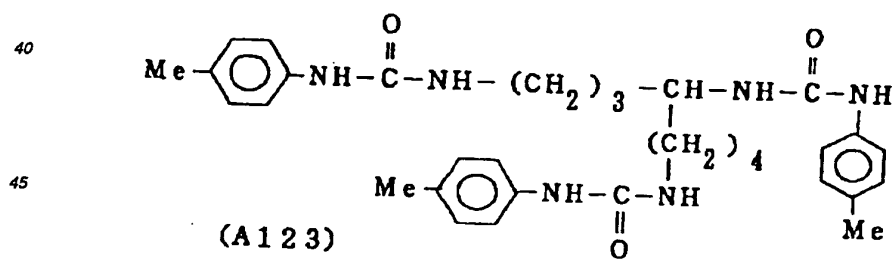
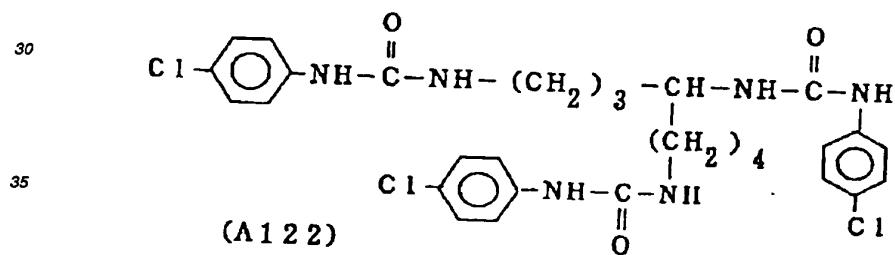
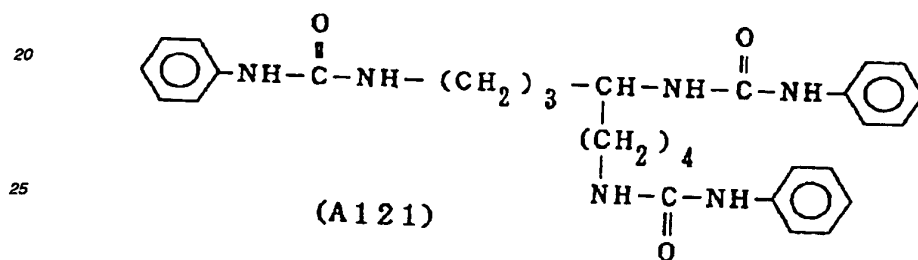
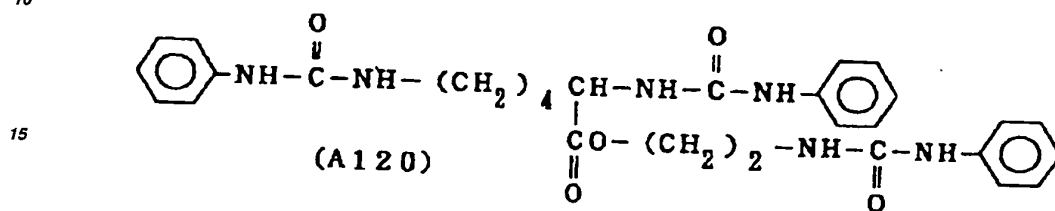
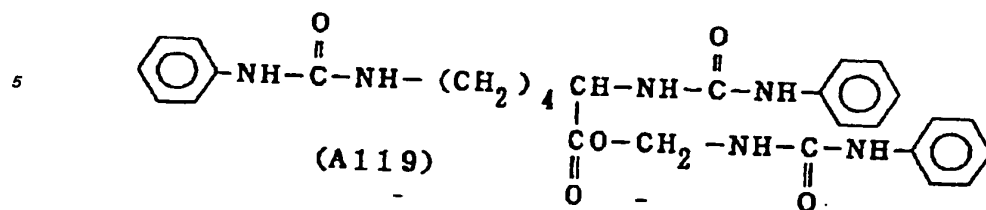


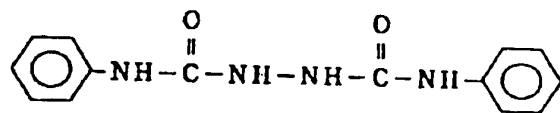
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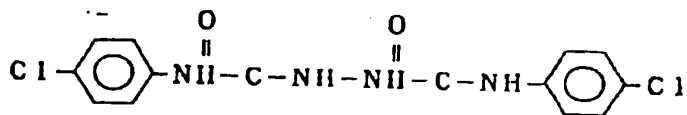




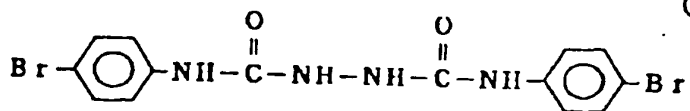




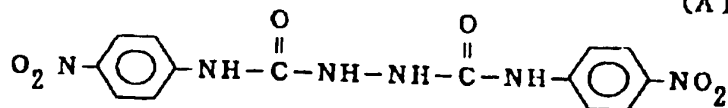
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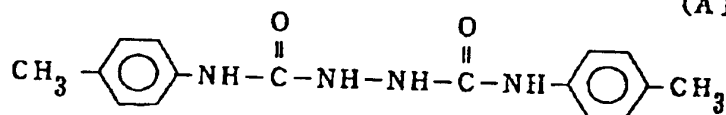
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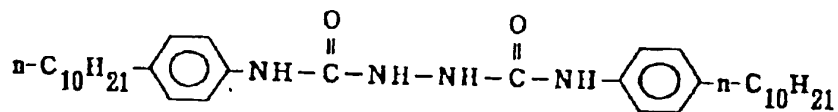
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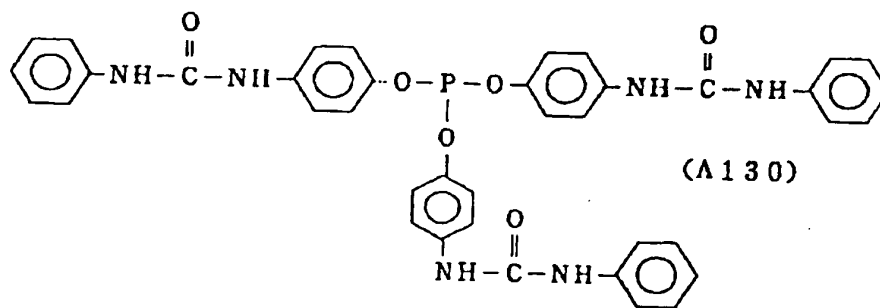
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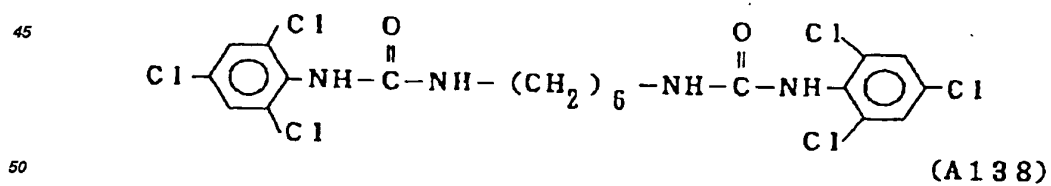
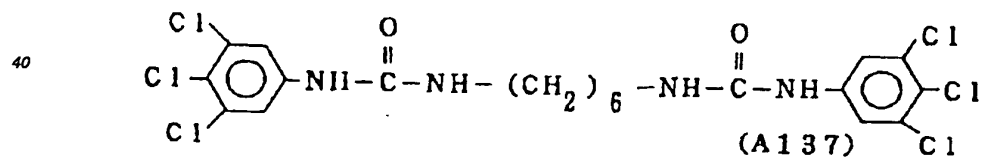
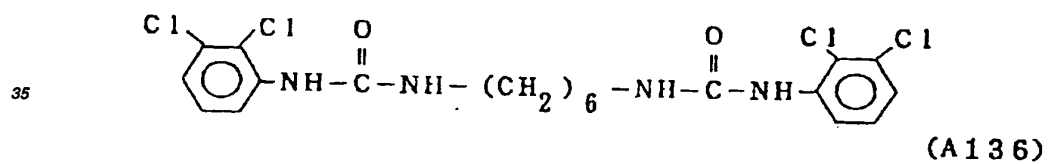
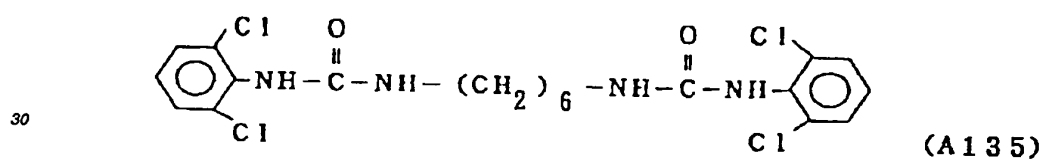
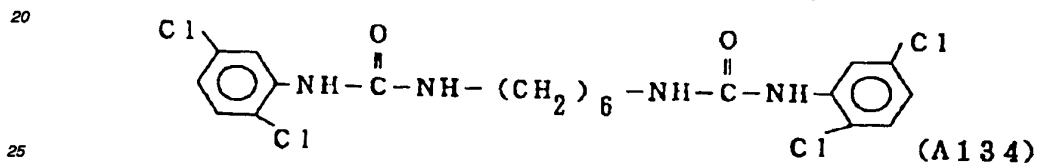
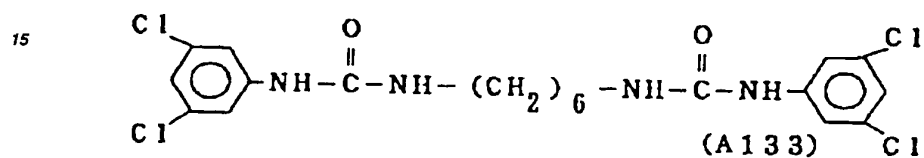
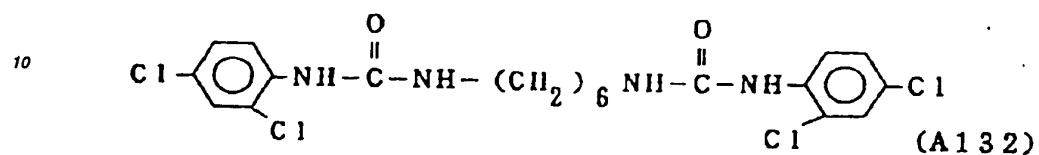
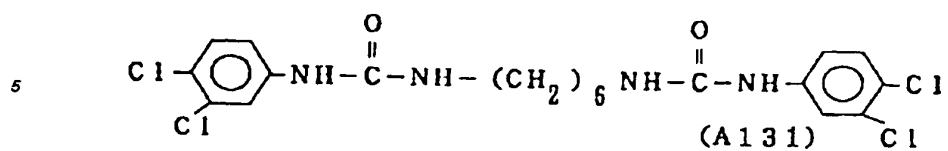
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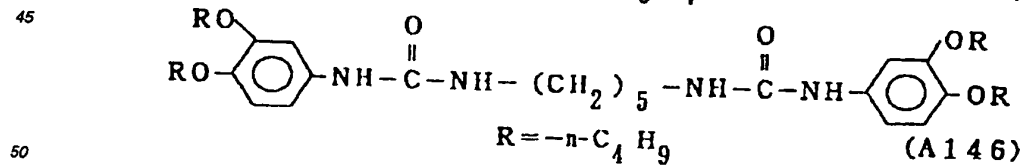
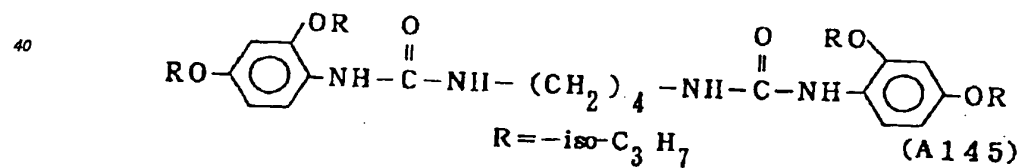
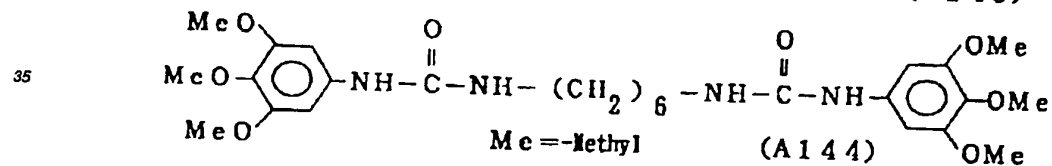
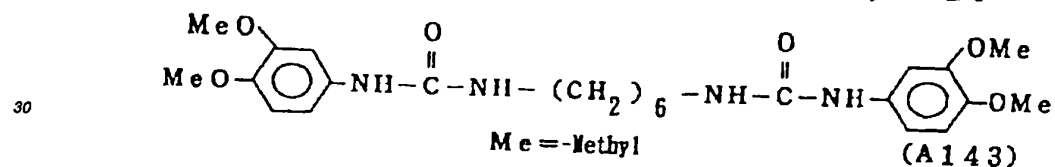
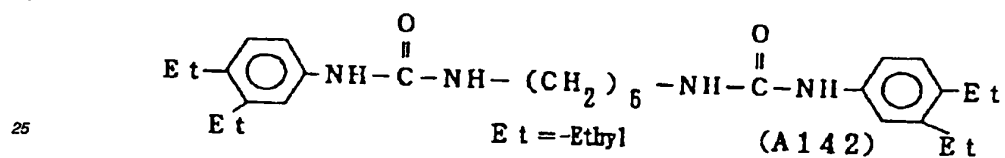
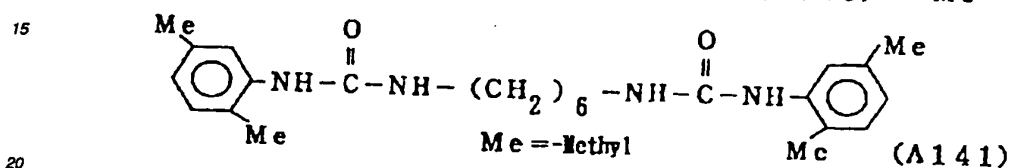
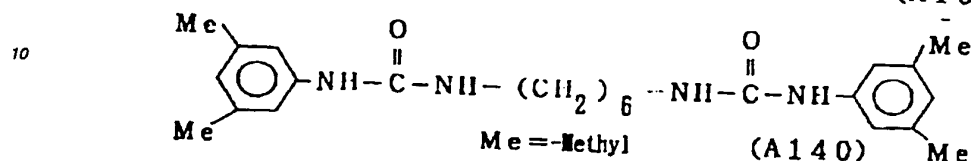
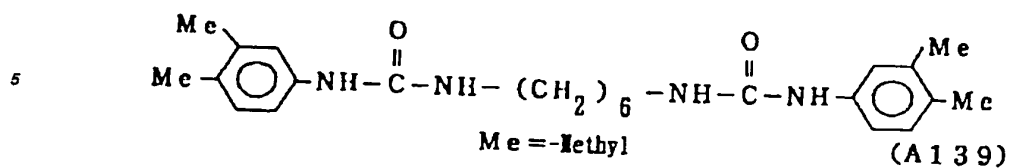


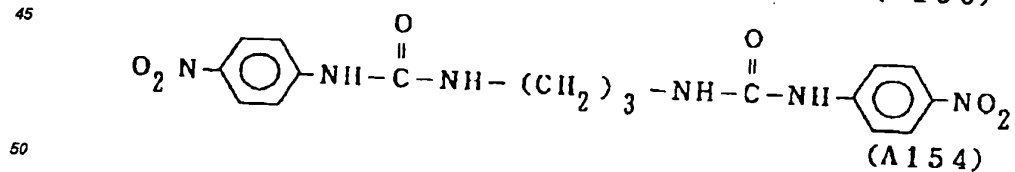
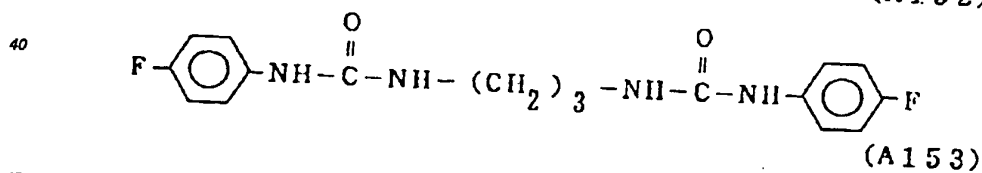
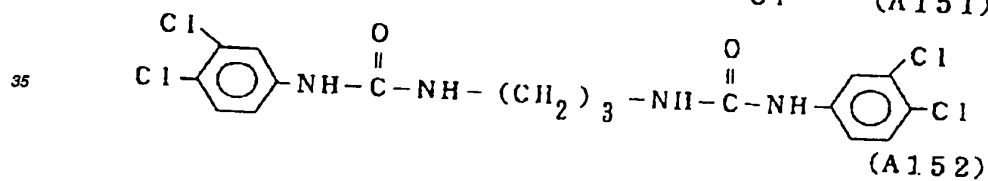
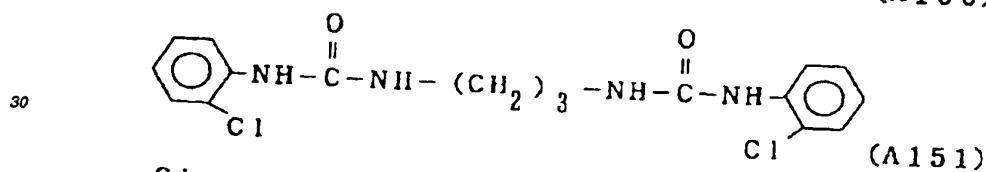
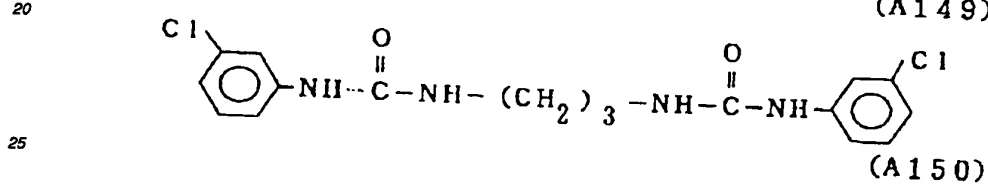
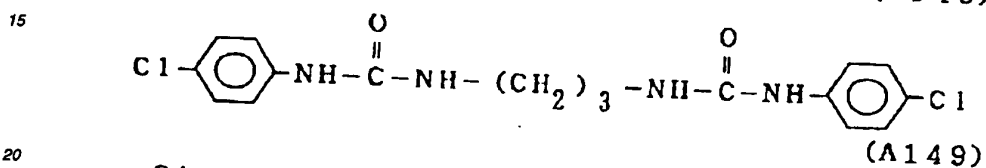
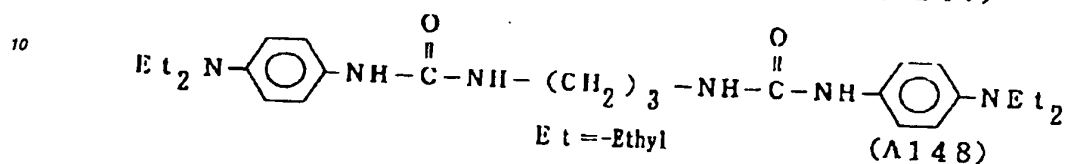
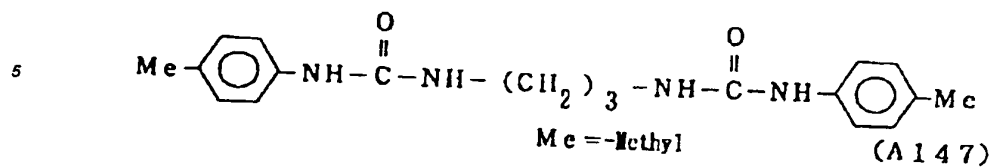
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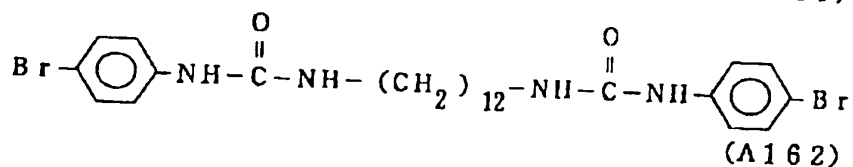
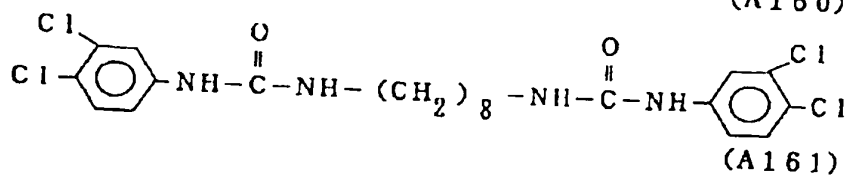
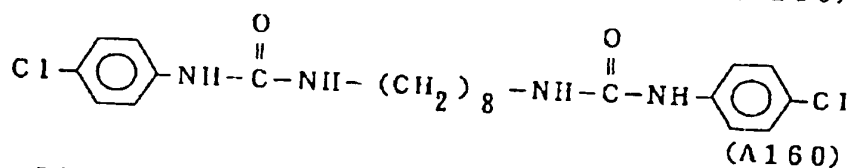
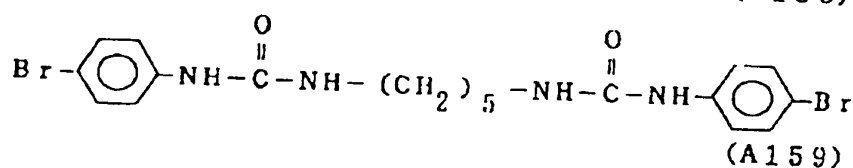
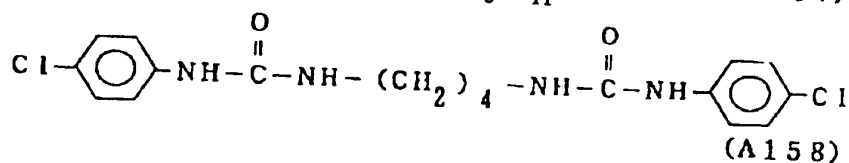
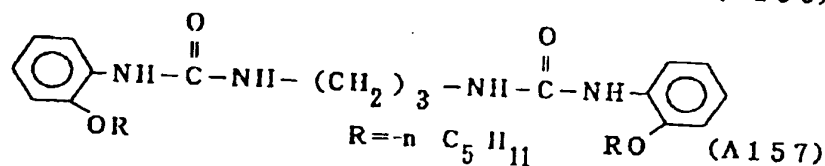
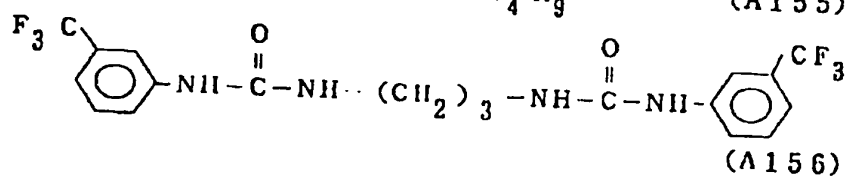
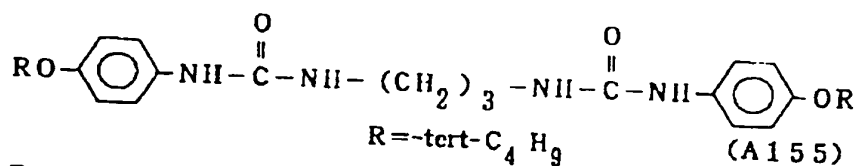


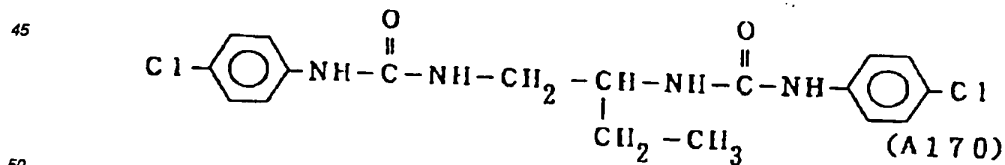
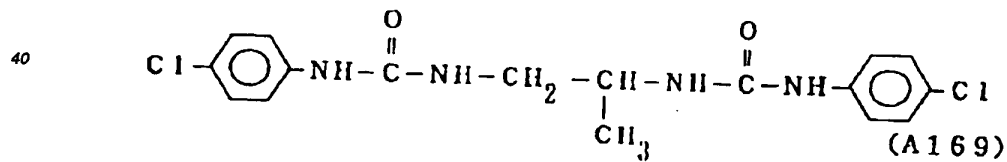
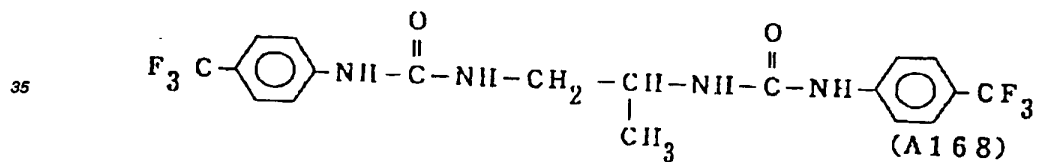
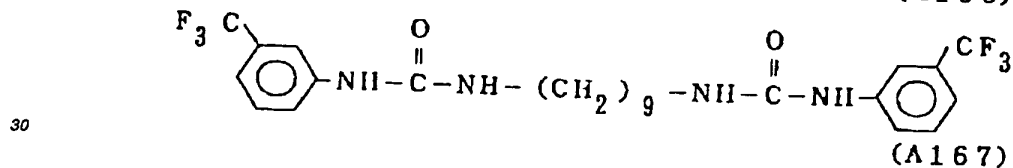
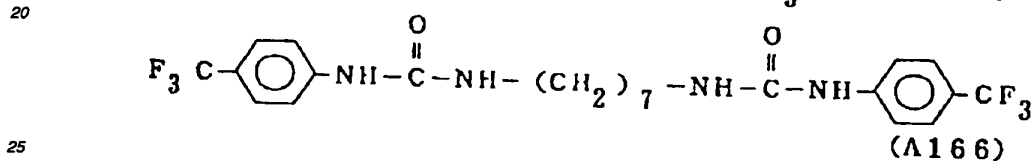
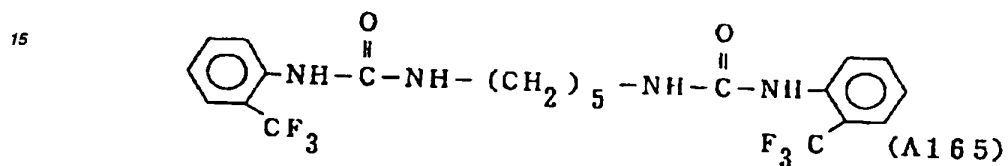
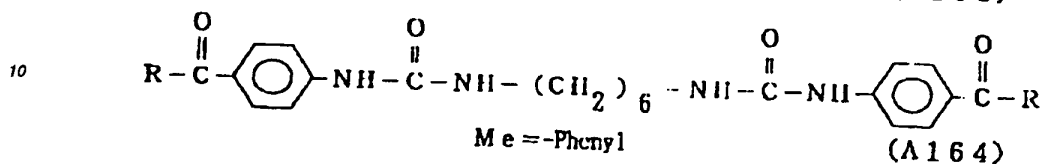
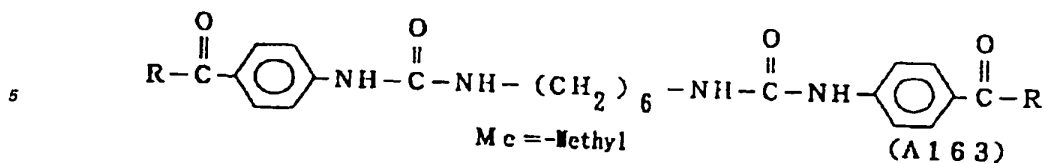
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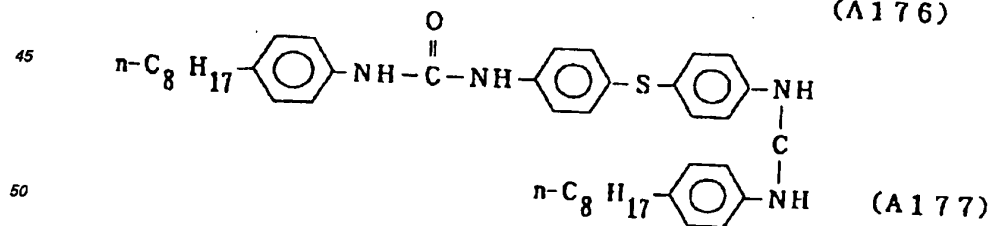
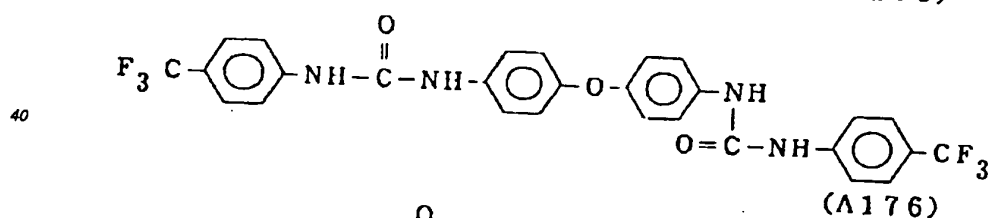
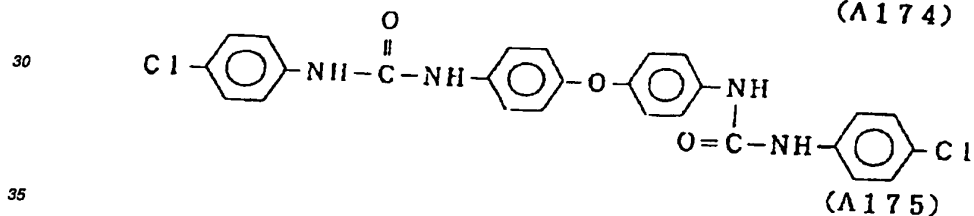
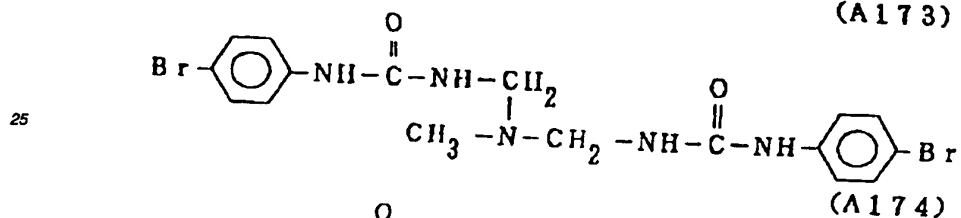
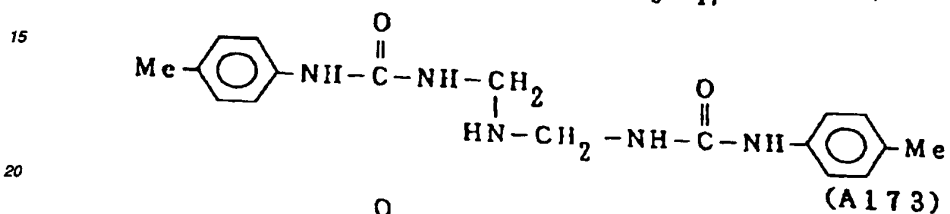
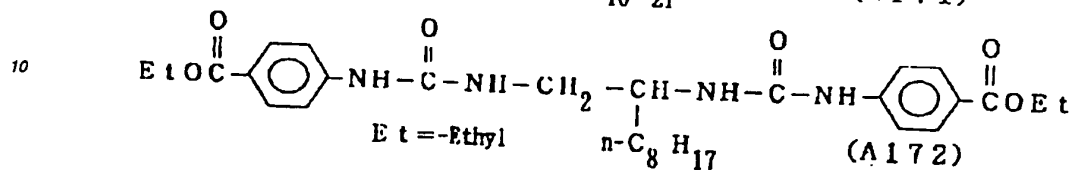
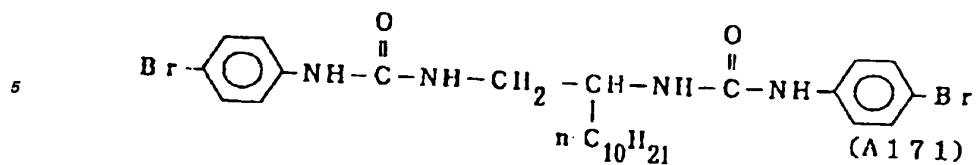


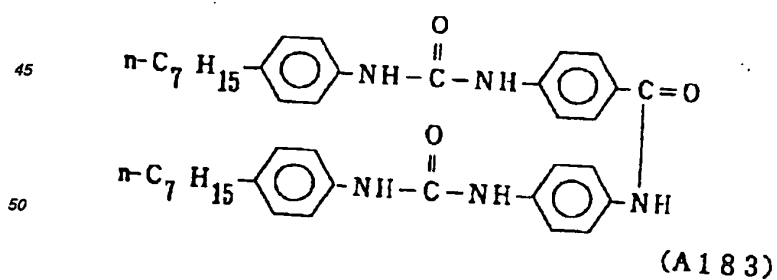
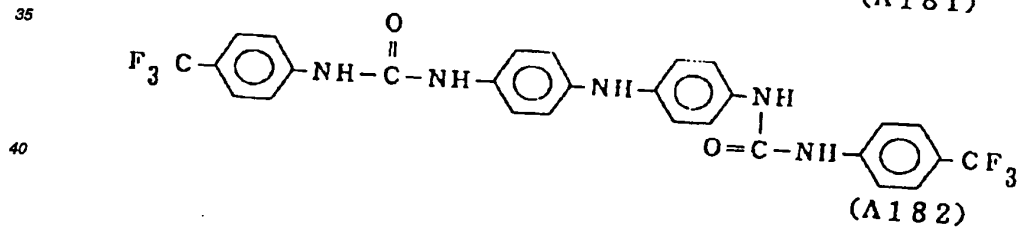
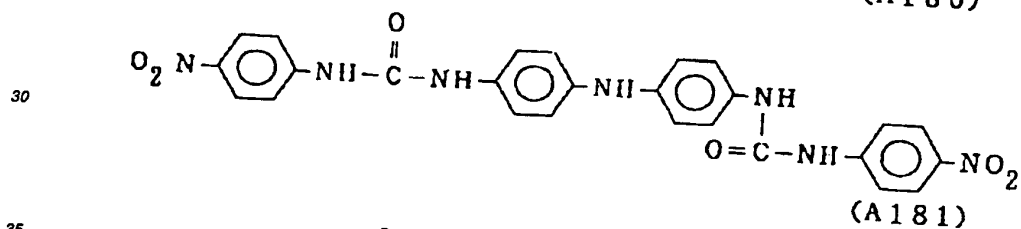
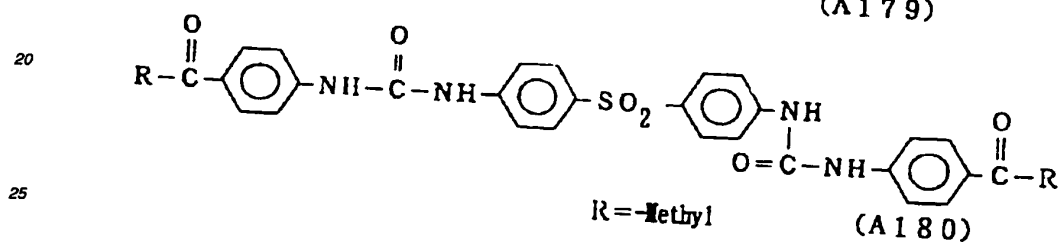
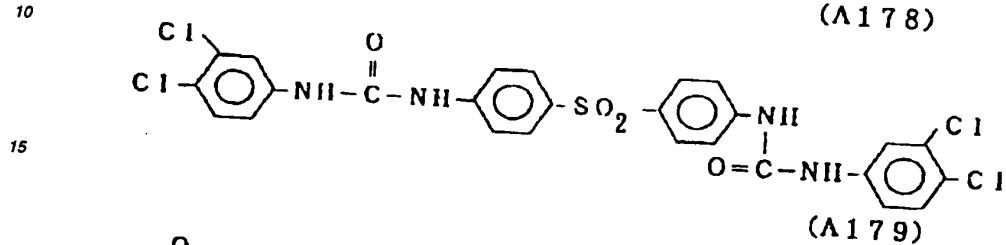
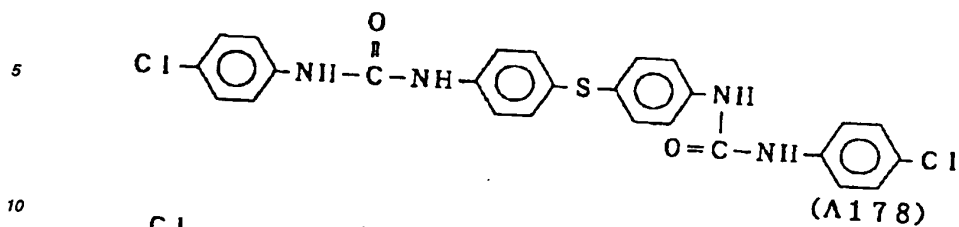


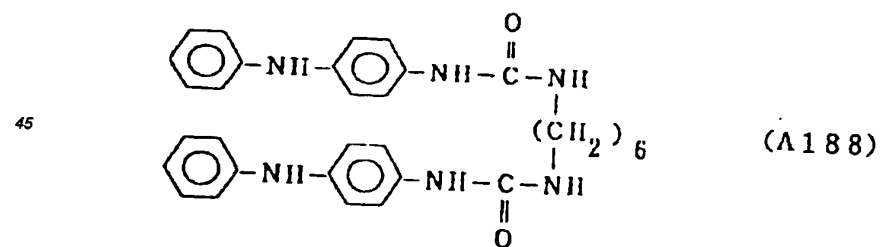
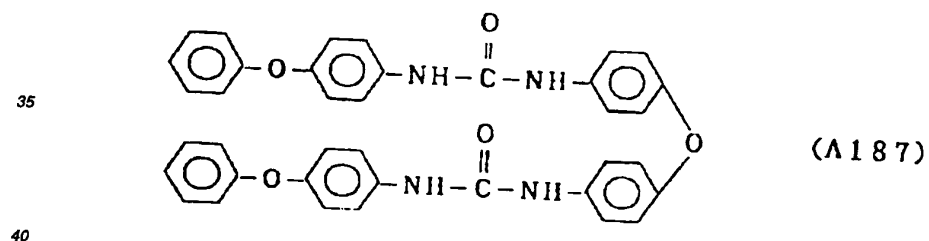
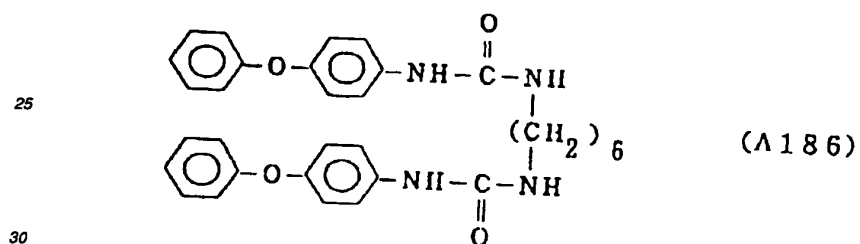
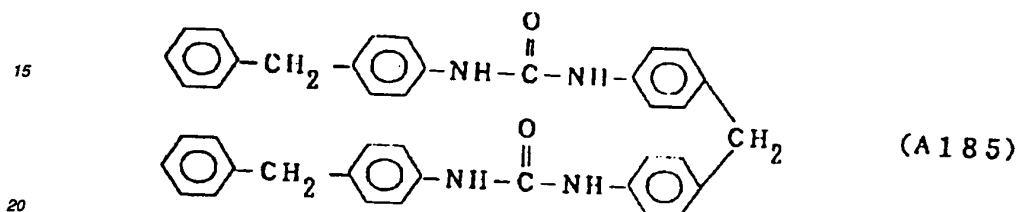
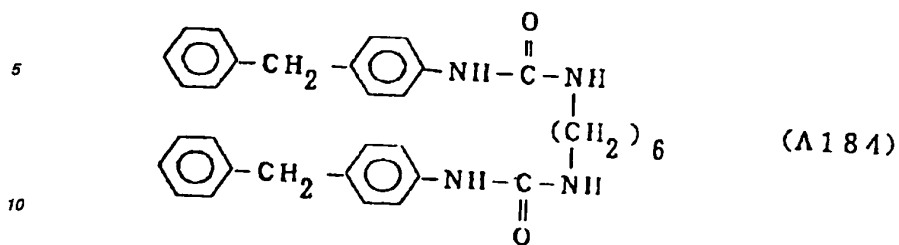


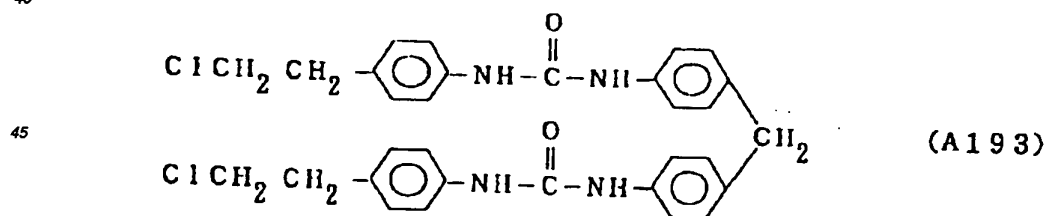
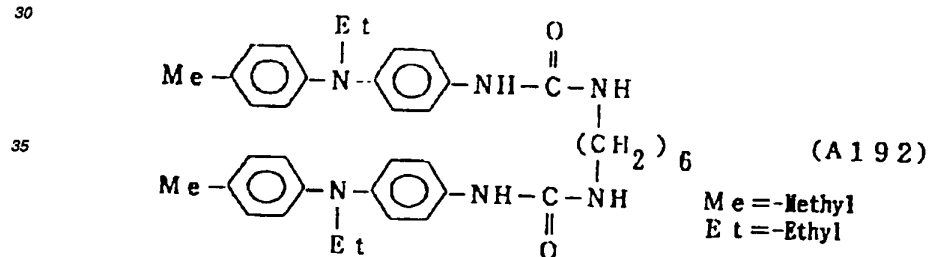
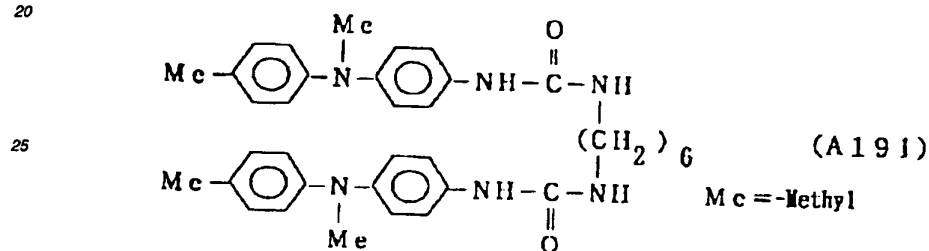
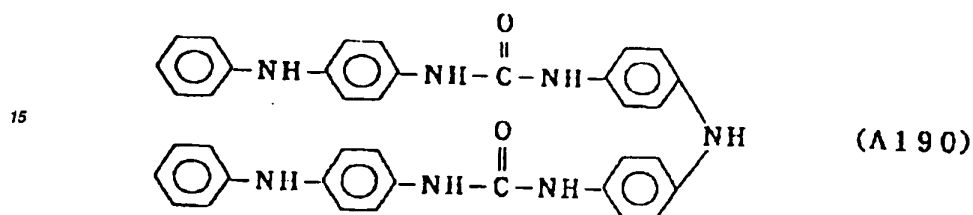
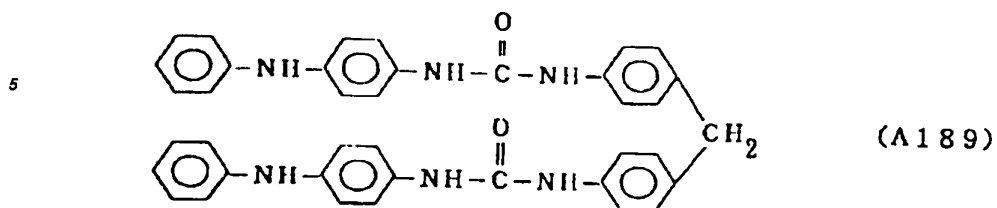
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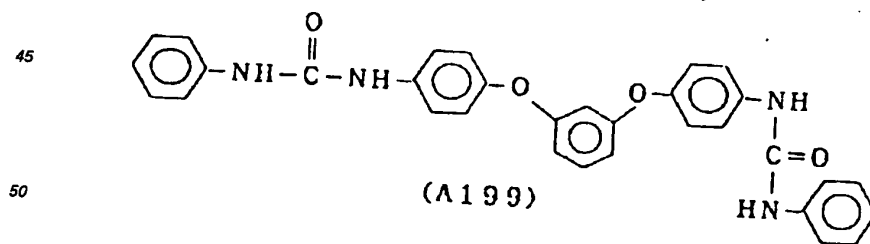
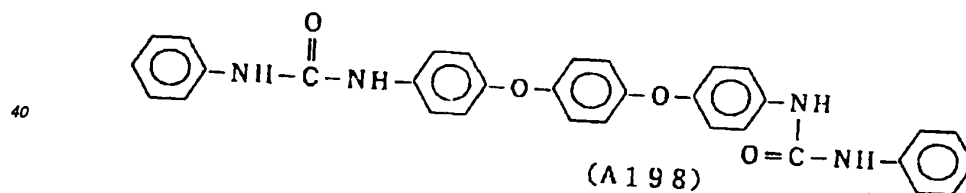
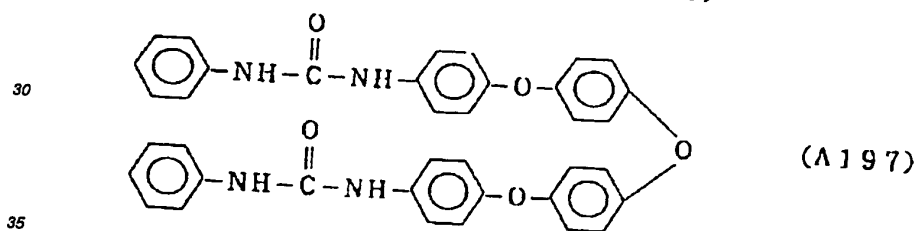
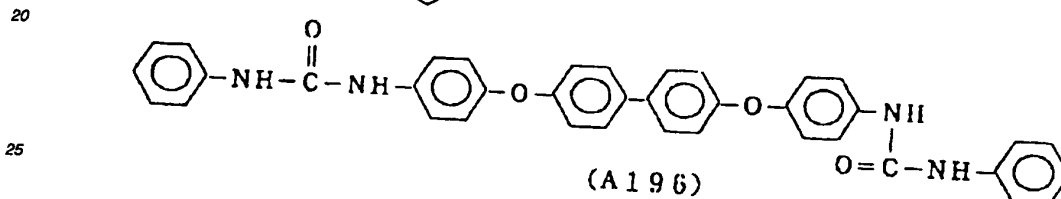
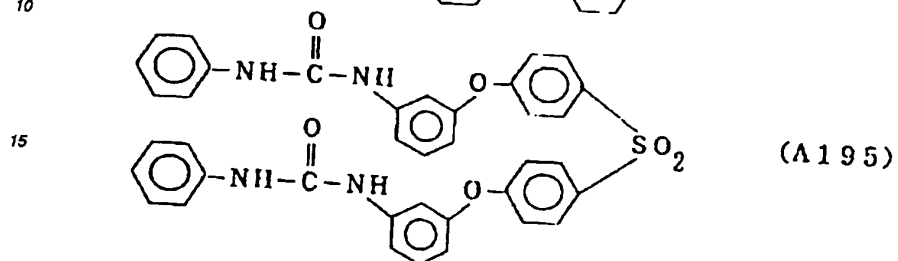
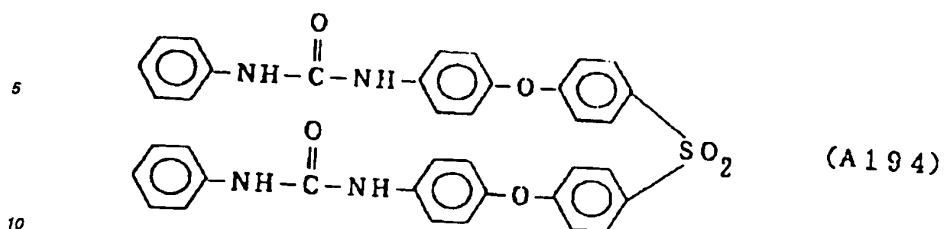
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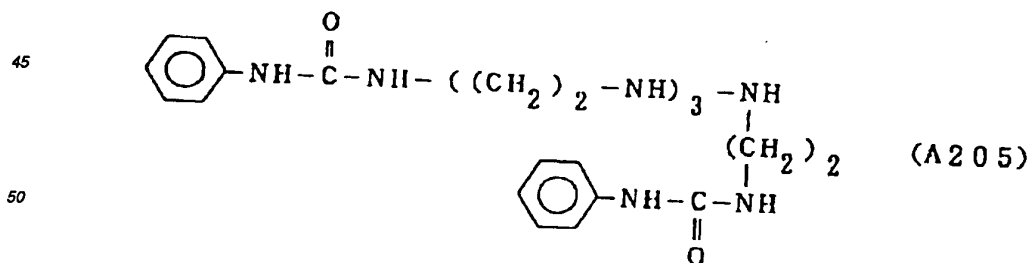
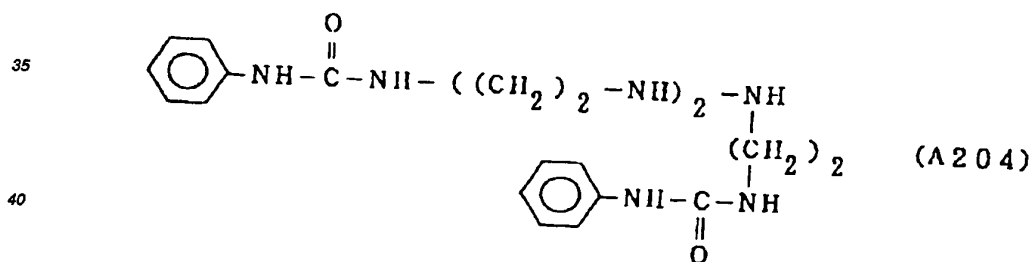
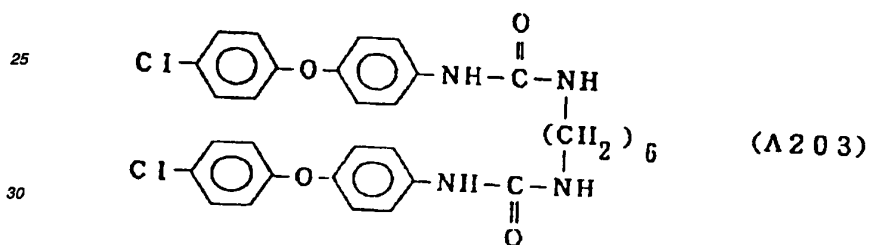
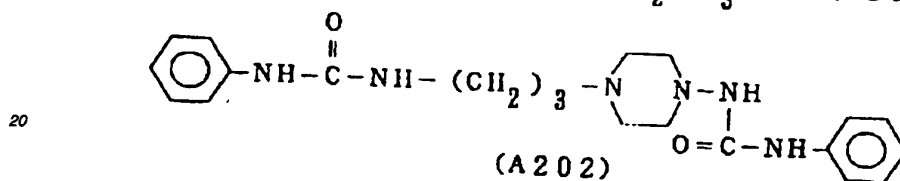
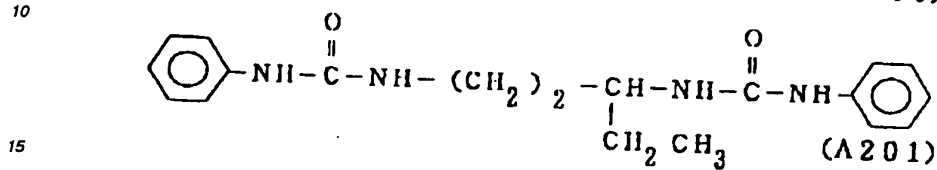
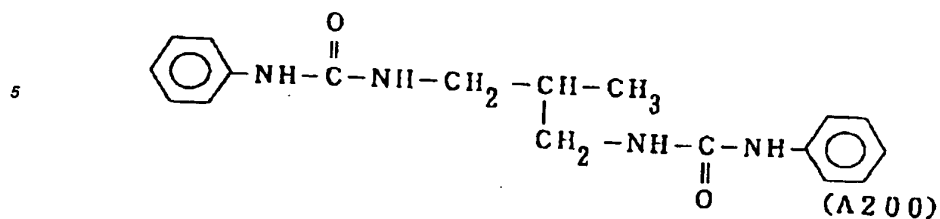


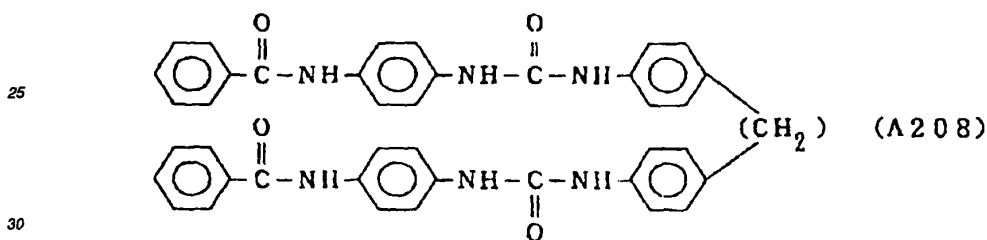
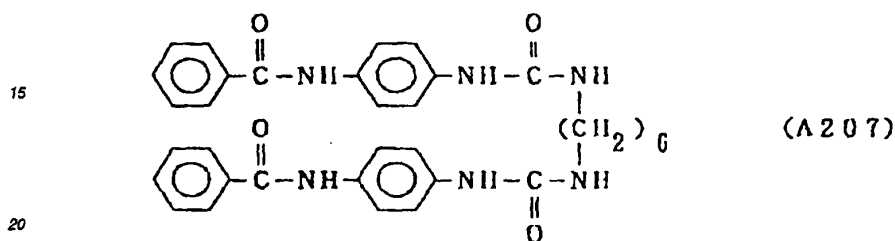
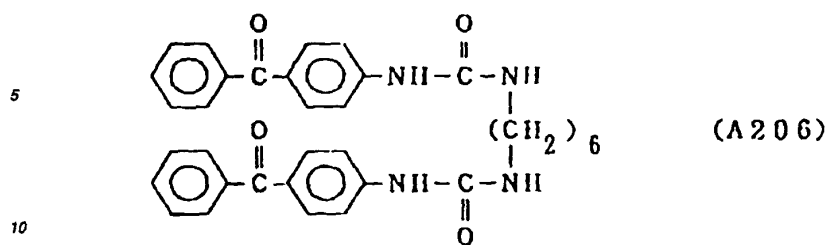






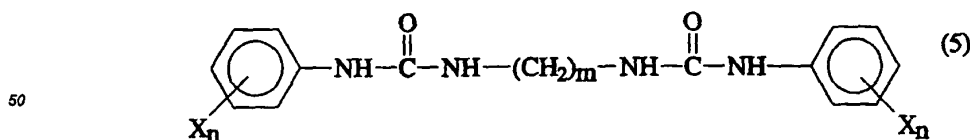






35 The divalent linking group is selected as appropriate according to properties such as the melting point, decomposition temperature and solubility in various solvents of the urea compound, and to the property of the thermal recording sheets using the urea compound as a developer. It is thus not specifically limited. For example, for the thermal recording sheet using the dimerized urea compound of Formula (3) as defined above, the divalent linking groups A are broadly divided as follows:

- 40 (1) A thermal recording sheet having a thermal recording layer containing at least one urea compound of formula (3), e.g. A1 - A12, A31 - A62, wherein the divalent group is a straight-chain alkylene group having 1 to 12 carbon atoms. In one aspect of the invention, therefore, the thermal recording sheet comprises as colour developer a urea compound of formula (5):



55 wherein X is as defined above in formula (1), n is an integer from 1 to 3, and m is an integer from 1 to 12.

(2) A thermal recording sheet wherein the colour developing is a compound of formula (3) as defined above, e.g. A13-A18, wherein divalent group (A) is a branched chain alkylene group having 1 to 15 carbon atoms.

(3) A thermal recording sheet wherein the colour developer is a compound of formula (3) as defined above, e.g.

A19-A23, A27-A30, wherein the divalent group A comprises two or more alkylene groups having 1 to 12 carbon atoms, the alkylene groups being linked to each other via nitrogen or oxygen.

Examples of alkylene groups linked to each other via nitrogen or oxygen include the following structures:

- 5 (Linked with nitrogen)
 $\text{-C(CH}_2\text{)}_{L1}\text{-NH-(CH}_2\text{)}_{M1}\text{-}$
 wherein L1 is an integer from 1 to 11, M1 is an integer from 1 to 11, and $L1 + M1 = 2$ to 12 .
 $\text{-(CH}_2\text{)}_{L2}\text{-NH-(CH}_2\text{)}_{M2}\text{-NH-(CH}_2\text{)}_{N2}\text{-}$
 wherein L2 is an integer from 1 to 10, M2 is an integer from 1 to 10, N2 is an integer from 1 to 10, and
 10 $L2 + M2 + N2 = 3$ to 12 .
 $\text{-(CH}_2\text{)}_{L3}\text{-NH-(CH}_2\text{)}_{M3}\text{-NH-(CH}_2\text{)}_{N3}\text{-NH-(CH}_2\text{)}_{Q3}\text{-}$
 wherein L3 is an integer from 1 to 9, M3 is an integer from 1 to 9, N3 is an integer from 1 to 9, Q3 is an integer from 1 to 9, and $L3 + M3 + Q3 = 4$ to 12 .
 $\text{-(CH}_2\text{)}_{L4}\text{-(NH-(CH}_2\text{)}_{M4}\text{)}_3\text{-NH-(CH}_2\text{)}_{N4}\text{-}$
 15 wherein L4 is an integer from 1 to 8, M4 is an integer from 1 to 2, N4 is an integer from 1 to 8, and $L4 + 3 \times M4 + N4 = 5$ to 12 .
 $\text{-(CH}_2\text{)}_{L5}\text{-(NH-(CH}_2\text{)}_{M5}\text{)}_4\text{-NH-(CH}_2\text{)}_{N5}\text{-}$
 wherein L5 is an integer from 1 to 7, M5 is an integer from 1 to 2, N5 is an integer from 1 to 7, and $L5 + 4 \times M5 + N5 = 6$ to 12 .
 20 (Linked with oxygen)
 $\text{-(CH}_2\text{)}_{L6}\text{-O-(CH}_2\text{)}_{M6}\text{-}$
 wherein L6 is an integer from 1 to 11, M6 is an integer from 1 to 11, and $L6 + M6 = 2$ to 12 .
 $\text{-(CH}_2\text{)}_{L7}\text{-O-(CH}_2\text{)}_{M7}\text{-O-(CH}_2\text{)}_{N7}\text{-}$
 wherein L7 is an integer from 1 to 10, M7 is an integer from 1 to 10, N7 is an integer from 1 to 10, and
 25 $L7 + M7 + N7 = 3$ to 12 .
 $\text{-(CH}_2\text{)}_{L8}\text{-O-(CH}_2\text{)}_{M8}\text{-O-(CH}_2\text{)}_{N8}\text{-O-(CH}_2\text{)}_{Q8}\text{-}$
 wherein L8 is an integer from 1 to 9, M8 is an integer from 1 to 9, N8 is an integer from 1 to 9, Q8 is an integer from 1 to 9, and $L8 + M8 + N8 + Q8 = 4$ to 12 .

30 When linked with nitrogen, hydrogen linked to the nitrogen atom of the joint group may be substituted with an alkyl group of 1 to 12 carbon atoms or an aryl group of 6 to 18 carbon atoms or unsubstituted.

(4) A thermal recording sheet having a thermal recording layer wherein the colour developer is a compound of formula (3), e.g. A103, A104, A114 or A115, wherein the divalent group A is a substituted or unsubstituted cycloalkyl group having 3 to 20 carbon atoms.

35 This category of colour developers is further divided into one in which the two phenylurea structures are linked directly to a single cycloalkyl ring, one in which the two phenylurea structures are linked to two cycloalkyls, and one in which the two phenylurea structures are linked to an alkylene group bonded to a cycloalkyl ring.

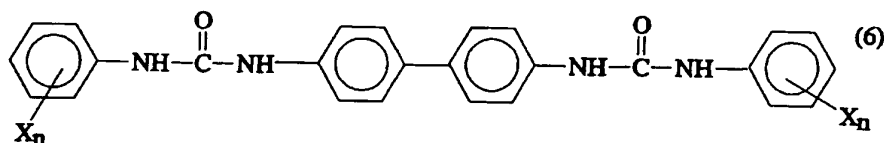
(5) A thermal recording sheet wherein the colour developer is a compound of formula (3), e.g. A87-A102, A107 or A108, wherein the divalent group A is a group having one substituted or unsubstituted aromatic ring having 6 to 20 carbon atoms.

40 This category of colour developers is further divided into one in which the two phenylurea structures are linked directly to one aromatic ring, and one in which the two phenylurea structures are linked to an alkylene group bonded to an aromatic ring. The aromatic ring can be, for instance, benzene, pyridine or naphthalene. Benzene is preferred in view of the availability of the raw material.

45 (6) A thermal recording sheet wherein the colour developer is a compound of formula (3), e.g. A63-A86 or A197 - A199, wherein the divalent group A is a group having at least two aromatic rings, each of which may be substituted or unsubstituted, which group has from 12 to 30 carbon atoms.

This category of colour developers is further divided into those having two aromatic rings, three aromatic rings, and four aromatic rings. Also in this case, benzene ring is preferable in view of availability of the raw material.

50 However, of the urea compounds of this category, the urea compound of Formula (6), that is, the divalent group A is diphenylene, is favoured for general applications in view of safety of the raw materials for synthesis.



wherein X is as defined above for formula (1), and n is an integer from 1 to 3.

(7) A thermal recording sheet wherein the colour developer is a compound of formula (3), e.g. A109-A111, wherein the divalent group A contains a substituted or unsubstituted piperazine ring and has from 4 to 12 carbon atoms.

In terms of the recording density, preferred options for the divalent group A are a straight-chain alkylene group having 1 to 12 carbon atoms, an alkylene group having 1 to 15 carbon atoms and having a branched chain, and two or more alkylene groups having 1 to 12 carbon atoms linked to each other via a nitrogen or oxygen atom. As regards the total balance of the recording density and heat resistance, preferred options for the divalent group A are a straight-chain alkylene group having 1 to 12 carbon atoms, and an alkylene group having 1 to 15 carbon atoms.

Further, the substituent (X) of the phenylurea structure of Formula (1) is also to be selected according to the melting point, decomposition temperature, the ease of handling in synthesis, and solubility in solvents of the urea compound, and the property of the thermal recording materials using the urea compound (recording density, heat resistance, and so on), as for the divalent group, and is not specifically limited. For instance, in terms of recording density, preferred options for X include hydrogen (that is, unsubstituted) and an electrophilic group (e.g. halogenated alkyl group, nitro group, cyano group, halogen atom). And in terms of the cost and safety of the urea compound, a preferred option for X is hydrogen.

On the other hand, a thermal recording sheet of the present invention which comprises a dimerized urea compound of Formula (2) or (3) or a trimerized urea compound of Formula (4) is superior in ground colour stability to heat and solvents. Even when the thermal recording sheet is placed in a high-temperature environment of above 120°C, the ground colour of the recording surface remains substantially unchanged. For example, the exemplified compound A8 does not develop a colour even at 150°C which is the highest temperature of static colour development. However, when the compound is heated by a thermal head to 200 to 300°C, it develops a dense colour. On the basis of common general knowledge of thermal recording sheets it would not be considered possible that no colour develops when the surface of the thermal recording sheet is contacted with a heat block at above 120°C, but can be recorded with a usable density by a thermal head. Such a material has been unknown.

Since the thermal recording sheet of the present invention has the above-described high heat resistance, it can be heat laminated with a plastic film on the recording surface after recording, used as an electrophotographic transfer sheet to be thermally fixed by coating a toner on the surface of the recording layer, and thermally fixed by coating a toner on the recording surface of the recorded sheet. Accordingly, the invention further provides a thermal recording card which comprises a thermal recording sheet of the invention as defined above, the said sheet being laminated with a plastic film. It also provides an electrophotographic transfer sheet which comprises a thermal recording sheet of the invention as defined above.

Further, the process for producing the thermal recording sheet of the present invention is very easy to control. In the production of conventional thermal recording sheets, the drying process after coating the recording layer requires very strict temperature control to prevent the coated surface from colour development, which limits high-speed coating. However, since the thermal recording sheet of the present invention does not develop ground colour even when contacted with hot air at 120°C, it can be dried at a high temperature. The control range of the drying temperature can therefore be substantially expanded, which leads to an improvement in productivity.

The thermal recording sheet of the present invention undergoes almost no change in ground colour even when contacted with solvents other than alcohol. This may be due to very low solubility in organic solvents of the dimerized or trimerized urea compound used. Therefore, it can be recorded by oil ink on the recording surface.

The inventors have conducted studies on the thermal decomposition temperature and solubility in various solvents of a number of dimerized or trimerized urea compounds, and have found that the thermal decomposition, temperature and solubility in various solvents can be controlled by suitable selection of the divalent linking group. By further studies, they have devised a recording sheet in which the recorded image can be erased by applying heat via a heat roll, a thermal head, a drying oven or a hot stamp, by irradiation with light by a laser or a halogen lamp, or by contacting with alcoholic solvents such as methanol or ethanol, and wherein the erased surface can be recorded again by means of a thermal head or a laser. This is a reversible recording sheet. It is achieved by using at least one dimerized urea com-

pound of Formula (2) or (3), or a trimerized urea compound of Formula (4), as defined above, as a colour developer in the thermal recording sheet of the invention.

According, the invention further provides a method of reversibly recording an image on a substrate, which method comprises:

5 recording an image on a thermal recording sheet of the invention as defined above, or an optical recording sheet of the invention as hereinafter defined;
erasing the recorded image by applying, respectively, heat or light to the recorded portion; and
recording an image again on the said sheet.

10 The invention also provides a method of reversibly recording an image on a substrate, which method comprises:

recording an image on a thermal recording sheet of the invention as defined above or an optical recording sheet of the invention as hereinafter defined:
15 contacting the recorded portion with an alcoholic solvent to erase the recorded image; and
recording an image again on the said sheet.

A reversible recording sheet which is good in erasability by a heat roll uses a colour developer which is a urea compound of Formula (5) (e.g. A1-A12) wherein the divalent linking group is a straight-chain alkylene group having 1 to 12
20 carbon atoms.

The thermal recording sheet using these urea compounds as colour developer, after being colour developed by means such as a thermal head, can be erased by contacting with a heat roll at 100 to 200°C and the erased surface can be recorded again by means such as a thermal head. For example, good results are obtained with a recording sheet in which the colour developer is a compound selected from the above-exemplified urea compounds A3, A4, A5,
25 A6, A8, A10, A14, and A49.

The dimerized urea compound of Formula (3) and the trimerized urea compound of Formula (4), from another point of view, are considered as compounds in which the phenylurea structures are appropriately separated by the divalent linking group. From this point of view, a urea compound which is superior in erasing ground colour under a heating condition different from recording by a thermal head is preferably a dimerized urea compound in which the phenylurea structures are separated by 3 to 8 carbon atoms.
30

The dimerized or trimerized urea compound used in the present invention alone in itself has a high colour developing ability as a colour developer in the thermal recording sheet, and the surface of the substrate coated with the compound along with the electron-donating dye precursor does not substantially develop a colour even at 120°C. It thus shows properties of superior resistance of ground colour to solvents. In addition, depending on the selected divalent linking group, some types of the urea compound of the present invention have a reversible recording function, as described above.
35

In a general method for producing the thermal recording sheet of the present invention, (a) the dye precursor and (b) the dimerized or trimerized urea compound as a colour developer are individually dispersed along with a binder having a dispersing function. As necessary, they are mixed with additives such as fillers, slip agents to obtain a coating colour, which is coated on a substrate by a conventional method known in the art, and then dried. The dimerized or trimerized urea compounds of the present invention can be used alone or in combination.
40

The dye precursor used in the thermal recording sheet of the present invention can be those which are known to the public in the area of thermal recording, and is not specifically limited, but triphenylmethane type leuco dyes, fluorane type leuco dyes, fluorene type leuco dyes, and the like are preferable. Typical dye precursors are shown below:

45 3,3-Bis(4'-dimethylaminophenyl)-6-dimethylaminophthalide
[alias Crystal Violet Lactone (CVL)]
3,3-Bis(4'-dimethylaminophenyl)-6-pyrrolidylphthalide
3,3-Bis(4'-dimethylaminophenyl)phthalide
50 [alias Malachite Green Lactone (MGL)]
Tris[4-(dimethylamino)phenyl]methane
[alias Leuco Crystal Violet (LCV)]
3-Dimethylamino-6-methyl-7-(m-trifluoromethylanilino) fluorane
3-Diethylamino-6-methyl-fluorane
55 3-Diethylamino-7-methyl-fluorane
3-Diethylamino-7-chlorofluorane
3-Diethylamino-6-methyl-7-chlorofluorane
3-Diethylamino-6-methyl-7-anilino fluorane

- 3-Diethylamino-6-methyl-7-p-methylanilino fluorane
 3-Diethylamino-6-methyl-7-(o,p-dimethylanilino) fluorane
 3-Diethylamino-6-methyl-7-(m-trifluoromethylanilino) fluorane
 3-Diethylamino-6-methyl-7-(o-chloroanilino) fluorane
 3-Diethylamino-6-methyl-7-(p-chloroanilino) fluorane
 3-Diethylamino-6-methyl-7-(o-fluoroanilino) fluorane
 3-Diethylamino-6-methyl-7-(p-n-butylanilino) fluorane
 3-Diethylamino-6-methyl-7-n-octylamino fluorane
 3-Diethylamino-6-chloro-7-anilino fluorane
 3-Diethylamino-6-ethoxyethyl-7-anilino fluorane
 3-Diethylamino-benzo[a] fluorane
 3-Diethylamino-benzo[c] fluorane
 3-Diethylamino-6-methyl-7-benzylamino fluorane
 3-Diethylamino-6-methyl-7-dibenzylamino fluorane
 3-Diethylamino-7-di(p-methylbenzyl)amino fluorane
 3-Diethylamino-6-methyl-7-diphenylmethylamino fluorane
 3-Diethylamino-7-dinaphthylmethylamino fluorane
 10-Diethylamino-4-dimethylaminobenzo[a] fluorane
 3-Dibutylamino-6-methyl fluorane
 3-Dibutylamino-6-methyl-7-chloro fluorane
 3-Dibutylamino-6-methyl-7-anilino fluorane
 3-Dibutylamino-6-methyl-7-p-methylanilino fluorane
 3-Dibutylamino-6-methyl-7-(o,p-dimethylanilino) fluorane
 3-Dibutylamino-6-methyl-7-(m-trifluoromethylanilino) fluorane
 3-Dibutylamino-6-methyl-7-(o-chloroanilino) fluorane
 3-Dibutylamino-6-methyl-7-(p-chloroanilino) fluorane
 3-Dibutylamino-6-methyl-7-(o-fluoroanilino) fluorane
 3-Dibutylamino-6-methyl-7-(p-n-butylanilino) fluorane
 3-Dibutylamino-6-methyl-7-n-octylamino fluorane
 3-Dibutylamino-6-chloro-7-anilino fluorane
 3-Dibutylamino-6-ethoxyethyl-7-anilino fluorane
 3-Di-n-pentylamino-6-methyl-7-anilino fluorane
 3-Di-n-pentylamino-6-methyl-7-(o,p-dimethylanilino) fluorane
 3-Di-n-pentylamino-6-methyl-7-(m-trifluoromethylanilino) fluorane
 3-Di-n-pentylamino-6-methyl-7-(o-chloroanilino) fluorane
 3-Di-n-pentylamino-6-methyl-7-(p-chloroanilino) fluorane
 3-Di-n-pentylamino-6-methyl-7-(o-fluoroanilino) fluorane
 3-Pyrrolidino-6-methyl-7-anilino fluorane
 3-Piperidino-6-methyl-7-anilino fluorane
 3-(N-methyl-N-n-propylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-n-propylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-isopropylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-n-butylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-isobutylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-n-hexylamino)-6-methyl-7-p-methylanilino fluorane
 3-(N-ethyl-N-n-hexylamino)-6-methyl-7-(o,p-dimethylanilino) fluorane
 3-(N-ethyl-N-n-hexylamino)-6-methyl-7-(m-trifluoromethylanilino) fluorane
 N3-(N-ethyl-N-n-hexylamino)-6-methyl-7-(o-chloroanilino) fluorane
 3-(N-ethyl-N-isoamylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-isoamylamino)-6-chloro-7-anilino fluorane
 3-(N-ethyl-N-3-methylbutylamino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-p-toluidino)-6-methyl-7-anilino fluorane
 3-(N-ethyl-N-p-toluidino)-6-methyl-7-(p-methylanilino) fluorane
 3-(N-ethyl-N-p-toluidino)-6-methyl-7-(o,p-dimethylanilino) fluorane
 3-(N-ethyl-N-tetrahydrofurfurylamino)-6-methyl-7-anilino fluorane
 3-(N-cyclohexyl-N-methylamino)-6-methyl-7-anilino fluorane
 3-(N-cyclohexyl-N-methylamino)-7-anilino fluorane
 3-(N-ethyl-N-3-methoxypropylamino)-6-methyl-7-anilino fluorane

- 3-(N-ethyl-N-3-ethoxypropylamino)-6-methyl-7-anilino fluorane
 2-(4-Oxaheptyl)-3-dimethylamino-6-methyl-7-anilino fluorane
 2-(4-Oxaheptyl)-3-diethylamino-6-methyl-7-anilino fluorane
 2-(4-Oxaheptyl)-3-dipropylamino-6-methyl-7-anilino fluorane
 5 3-(4"-Aminostilbuldy-4'-amino)-7,8-benzofuran
 3,6,6'-Tris(dimethylamino)spiro[fluorene-9,3'-phthalide]
 3,6,6'-Tris(diethylamino)spiro[fluorene-9,3'-phthalide]
 3-(4-Diethylamino-2-ethoxyphenyl)-3-(1-ethyl-2-methylindol-3-yl)-4-azaphthalide
 3-(4-Diethylamino-2-ethoxyphenyl)-3-(1-octyl-2-methylindol-3-yl)-4-azaphthalide
 10 3-(4-Diethylamino-2-n-hexylphenyl)-3-(1-ethyl-2-methylindol-3-yl)-4-azaphthalide
 3-(4-Cyclohexylmethylamino-2-methoxyphenyl)-3-(1-ethyl-2-methylindol-3-yl)-4-azaphthalide
 3-(4-Cyclohexylethylamino-2-methoxyphenyl)-3-(1-ethyl-2-methylindol-3-yl)-4-azaphthalide
 3,3-Bis(1-ethyl-2-methylindol-3-yl)phthalide
 3,3-Bis(1-methyl-2-octylindol-3-yl)phthalide
 15 3-(1-Ethyl-2-methylindol-3-yl)-3-(1-n-butyl-2-methylindol-3-yl)phthalide
 3,7-Bis(dimethylamino)-10-benzoylphenothiazine
 3,7-Bis(dimethylamino)-N-[p-N-bis(4,4'-dimethylaminophenyl)methylamino]benzoylphenothiazine
 3,7-Bis(dimethylamino)-N-[p-N-bis(4,4'-diethylaminophenyl)methyl]benzoylphenothiazine
 3,6-Bis(diethylamino)fluorane- γ -(2'-nitro)anilino lactam
 20 3,6-Bis(diethylamino)fluorane- γ -(3'-nitro)anilino lactam
 3,6-Bis(diethylamino)fluorane- γ -(4'-nitro)anilino lactam
 3,6-Bis(diethylamino)fluorane- γ -anilino lactam.

These dye precursors may be used alone or as mixtures of two or more types. Since fluorane type dye precursors
 25 are particularly high in stability of color development and ground color as basic functions of the thermal recording sheet,
 they can be preferably used in the present invention. Naturally, when thermal stability is important, a dye precursor
 which is high in melting point and decomposition temperature is preferable. When the reversible recording function is
 important, it is preferable to use 3-dimethylamino-7-(m-trifluoromethyl)anilino)fluorane and the like, which is weak in
 image stability, can withstand repeated use, and is high in decomposition temperature.

30 The binder usable in the present invention includes completely-hydrolyzed polyvinylalcohol having a polymerization
 degree of 200 to 1900, partially-hydrolyzed polyvinylalcohol, carboxy-modified polyvinylalcohol, amide-modified polyvi-
 nylalcohol, sulfonic acid-modified polyvinylalcohol, butyral-modified polyvinylalcohol, other modified polyvinylalcohols,
 styrene-maleic anhydride copolymer, styrene-butadiene copolymer, cellulose derivatives such as hydroxyethylcellu-
 35 lose, methylcellulose, carboxymethylcellulose, ethylcellulose and acetylcellulose, polyvinylchloride, polyvinylacetate,
 polyacrylamide, polyacrylic esters, polyvinylbutyral, polystyrene and copolymers thereof, polyamide resins, silicone res-
 ins, petroleum resins, terpene resins, ketone resins, and coumarone resins. Of these binders, the polyvinylalcohol type
 binder is preferable in terms of the dispersibility, binding properties, and thermal stability of ground color. For reversible
 recording, a binder which can withstand repeated use and is small in degradation is preferable. These binders are used
 40 by dissolving in solvents such as water, alcohol, ketone, ester, and hydrocarbon, emulsifying in water or other solvents,
 or dispersing to a paste form, and can be used in combination according to the quality requirements.

In the present invention, when a thermal recording sheet which is particularly high in thermal stability of ground
 color is produced, it is better in principle not to use a sensitizer. When a sensitizer is used, it tends to melt at a drying
 temperature, and causes the dye precursor to react with the color developer to develop the ground color. However, on
 the other hand, since the sensitizer promotes the erasing function, when importance is attached to the erasing function
 45 rather than the thermal stability of ground color, that is, the reversible recording function, a sensitizer may be used. Sen-
 sitizers used for this purpose include 2-di(3-methylphenoxy)ethane, p-benzylbiphenyl, β -benzyloxynaphthalene, phenyl
 1-hydroxy-2-naphthoate, dibenzylterephthalate, benzyl p-benzyloxybenzoate, diphenylcarbonate, ditolylcarbonate, 4-
 biphenyl-p-tolylether, m-terphenyl, 1,2-diphenoxyethane, 1,2-bis(m-tolyl)oxyethane, di(p-methylbenzyl)oxalate, and
 di(p-chlorobenzyl)oxalate.

50 The filler used in the present invention includes inorganic fillers such as silica, calcium carbonate, kaolin, calcined
 kaolin, diatomaceous earth, talc, zinc oxide, titanium oxide, zinc hydroxide, and aluminum hydroxide; polystyrene-
 based organic fillers, styrene/butadiene-based organic fillers, styrene/acrylic-based organic fillers, and hollow organic
 fillers.

In addition to the above, a release agent such as fatty acid metal salts, a slip agent such as waxes, benzophenone
 55 or benzotriazole type ultraviolet absorbers, a water-resistant agent such as glyoxal, a dispersant, a defoamer, and the
 like can be used.

Types and ratios of (a) the dye precursor, (2) the dimerized or trimerized urea compound, and other ingredients are
 determined by the required properties and recording adaptability, and are not specifically limited but, normally, based

on one part of the dye precursor, 1 to 8 parts of the dimerized or trimerized urea compound, and 1 to 20 parts of the filler are used, and the binder is used in an amount of 10 to 25 % by weight to the total solid. For repeated use such as the reversible recording sheet, it is preferable to use a composition as simple as possible. These materials are finely crushed by a crusher such as a ball mill, an attriter, or a sand grinder, or an appropriate emulsifying apparatus to a particle diameter of several microns or less, a binder and, as necessary, other additives are added to obtain a coating color. The coating color comprising the above composition is coated on a desired substrate such as paper, synthetic paper, non-woven fabrics, plastic films, plastic sheets, or composite sheets thereof to obtain the objective thermal recording sheet.

Further, an overcoating layer comprising a polymer can be provided on top of the recording layer to enhance the preservability, or an undercoating layer of a polymer containing a filler can be provided under the recording layer to enhance the color developing sensitivity.

The thermal recording sheet of the present invention, utilizing its high ground color stability, may be heat laminated with a plastic film to form a transparent and strong protective coating. For example, even after thermal recording, a heat-resistant card can be easily prepared using a commercial simple laminator.

Of the thermal recording sheets of the present invention, one which has an erasing function is useful as a reversible recording sheet (rewrite or rewritable recording sheet), or as a simple displaying sheet. However, for the latter application, it is necessary to achieve recording and erasing almost simultaneously.

Erasing of the thermal recording sheet of the present invention is achieved by two methods. One method uses a heat roll, a thermal head, a hot stamp, a carbon dioxide laser, a semiconductor laser, sunlight, a halogen lamp, or the like to erase the recorded image by heat. For example, for a heat roll, the erasing temperature is preferably 100 to 200°C with a feed speed of 8 to 45 mm/sec. The other method uses an alcoholic solvent to erase the recorded image.

The thermal recording sheet of the present invention may contain an optical absorbent which absorbs light to convert it to heat in the thermal recording layer. The present invention therefore also provides an optical recording sheet which comprises a thermal recording sheet of the invention as defined above and an optical absorbent. The optical absorbent can be a substance which absorbs wavelengths of various light sources, and is not specifically limited. The optical recording sheet may be laminated with a plastic film to form a thermal recording card, which is a further aspect of the invention. A yet further aspect of the invention is an electrophotographic transfer sheet which comprises an optical recording sheet of the invention as defined above.

For example, for a recording light source having continuous wavelength, such as a stroboflash, the optical absorbent can be a heat reaction product of thiourea derivative/copper compound described in JP-A-2-206583 and 5-30954, graphite, copper sulfide, molybdenum trisulfide, black titanium, and the like described in JP-A-3-86580, or carbon black.

On the other hand, when a semiconductor laser is used as a recording light source, the optical absorbent can be polymethine type dyes (cyanine dyes); azolenium type dyes, pyrylium type dyes, thiopyrylium type dyes, squarylium type dyes, chroconium type dyes, dithiol complexes, mercaptophenol-metal complex type dyes, mercaptanaphthol-metal complex type dyes, phthalocyanine type dyes, naphthalocyanine type dyes, triarylmethane type dyes, immonium type dyes, diimmonium type dyes, naphthoquinone type dyes, anthraquinone type dyes, and metal complex type dyes. Further, the optical absorbents listed for a light source having continuous wavelength can also be used as well.

Specifically, near infrared absorption dyes described in "Colour chemical dictionary" (The society of Synthetic Organic Chemistry, CMC-publishing company, 1988, P196-200) "Kagaku to Kogyo" (Vol 5. 1986, p376-389), JP-A-61-69991 and 61-246391, and US patents 3251881, 35570122, 3575871 and 3637769; 1,1,5,5-tetrakis(p-dimethylaminophenyl)-3-methoxy-1,4-pentadiene (or its cation form), 1,1,5,5-tetrakis(p-diethylaminophenyl)-3-methoxy-1,4-pentadiene (or its cation form); toluenedithiol nickel complex, 4-tert-butyl-1,2-benzenedithiol nickel complex, bisdithiobenzilnickelcomplex, bis(4-ethylthiobenzil)nickel complex, and so on. These optical absorbents can be used alone or as mixture of two or more types.

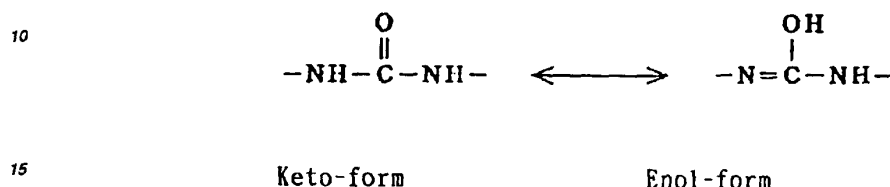
These optical absorbents may be used by: (a) a method in which the optical absorbent is simply mixed in the materials necessary for the thermal recording sheet, (b) a method in which the optical absorbent is previously melted and mixed, and dissolved or dispersed to be used, as described in Japanese OPI 2-217287, or (c) a method in which the optical absorbent is previously dissolved or dispersed by a solvent in the materials necessary for the thermal recording sheet, the solvent is removed from the mixture, and then used. The optical absorbent may also be co-dispersed with the color developer with the dye precursor, with the color developer and the sensitizer or with the dye precursor and the sensitizer.

The thermal recording sheet of the present invention, even when combined with the optical absorbent to form an optical recording sheet, is substantially unchanged in the basic properties (e.g. ground color stability such as heat resistance and solvent resistance). Even the optical recording sheet can be heat laminated or toner recorded. This is also true for the additional functions (reversible recording). However, when a sensitizer is used, the heat resistance tends to be impaired. The thermal recording sheet laminated with the plastic film, which contains the optical absorbent, can be recorded by light and as laser-light.

The novel dimerized or trimerized urea compound according to the present invention is a color developer which is

superior in color development and ground color stability to heat and solvents as basic functions, and some types have a reversible recording function. The reason for the superior ground color stability and the reversible recording function has yet to be clarified. However, this is considered as follows:

Depending on the condition, the dimerized or trimerized urea compound of the present invention changes in structure as shown below. Since this change is a phenomenon similar to the keto/enol tautomerism, the structures are referred to here as the keto-form and the enol-form for convenience.



It is considered that the dimerized or trimerized urea compound must have the enol-form to function as a color developer. Since enol-formation requires a high temperature, and the thermal head instantaneously provides a high temperature of 200 to 300°C, the urea compound contacting with the thermal head undergoes enol-formation to have a color developing function, opening the lactone ring of the dye precursor to develop a color. Therefore, the urea compound does not change until the enol-formation temperature is reached, does not react with the dye precursor, and the ground color remains unchanged. This would be the reason for the high heat resistance. Further, since the dimerized or trimerized urea compound increases the number of active hydrogens compared to a monourea compound, a good color developing function is obtained.

On the other hand, if the thus produced enol-form converts to the keto-form for some reason, erasing would occur. Therefore, when the compound is provided with an appropriate temperature and head, or contacts with an alcoholic solvent, keto-formation occurs, resulting in erasing. Since enol-formation and keto-formation occur under quite different conditions, the enol-form and the keto-form can be repeated under the individual conditions, thereby enabling reversible recording.

The reason why the ground color is not changed by writing with an oil ink is considered as due to the fact that the dimerized or trimerized urea compound of the present invention is very low in solubility in the solvent used in the oil ink, and the dye precursor and the color developer are not substantially mixed with each other even when contacting the solvent.

The invention will be further described in the Examples which follow.

Examples

Synthesis of dimerized and trimerized urea compounds

Dimerized and trimerized urea compounds were synthesized by the following synthetic methods. General synthetic methods are shown below.

Synthesis Example 1

A monoisocyanate compound (2.2 eq.) was dissolved in ethyl acetate (or acetone). Into the solution, a solution of a diamine compound (1.0 eq.) in ethyl acetate (or acetone) was dropped. After stirring for a predetermined time, generated precipitate was filtered, and washed with ethyl acetate, n-hexane, hot water, and methanol in this order until 1 spot was obtained on TLC (thin layer chromatograph) to obtain a urea compound.

Synthesis Example 2

A diisocyanate compound (1.0 eq.) was dissolved in ethyl acetate (or acetone). Into the solution, a solution of a monoamine compound (2.2 eq.) in ethyl acetate (or acetone) was dropped. After stirring for a predetermined time, generated precipitate was filtered, and washed with ethyl acetate, n-hexane, hot water, and methanol in this order until 1 spot was obtained on TLC to obtain a urea compound.

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Synthesis Example 3

A triisocyanate compound (1.0 eq.) was dissolved in ethyl acetate (or acetone). Into the solution, a solution of a monoamine compound (3.3 eq.) in ethyl acetate (or acetone) was dropped. After stirring for a predetermined time, generated precipitate was filtered, and washed with ethyl acetate, n-hexane, hot water, and methanol in this order until 1 spot was obtained on TLC to obtain a urea compound.

Production of thermal recording sheet

In the following description, unless otherwise noted, part and % indicate part by weight and % by weight, respectively.

Examples 1-48

As shown below, thermal recording sheets were produced using 3-N,N-diethylamino-6-methyl-7-anilino-fluorane (ODB) as a dye precursor and the dimerized urea compound (or trimerized urea compound) of the present invention (Table 1, Table 2, Table 3) as a developer.

Specifically, a color developer dispersion (Solution A) and a dye precursor dispersion (Solution B) of the following compositions were milled by a sand grinder to an average particle diameter of 1 micron.

(Solution A: color developer dispersion)

Inventive dimerized urea compound (or trimerized urea compound)	6.0 parts
10% Aqueous polyvinylalcohol solution	18.8
Water	11.2

(Solution B: dye precursor dispersion)

3-N,N-diethylamino-6-methyl-7-anilino-fluorane (ODB)	2.0 parts
10% Aqueous polyvinylalcohol solution	4.6
Water	2.6

Then, the Solution A (color developer dispersion), the Solution B (dye precursor dispersion), and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

Solution A (color developer dispersion)	36.0 parts
Solution B (dye precursor dispersion)	9.2
Kaolin clay (50% dispersion)	12.0

The coating color was coated on one side of a 50 g/m² base paper to a coating coverage of 6.0 g/m², dried, and supercalendered to a flatness of 500 to 60 seconds to obtain a thermal recording sheet.

Examples 49-56

Thermal recording sheets were produced using the dye precursors other than ODB and using the same procedure as in Examples 1-48 (Table 5).

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(Dye precursor)

ODB-2: 3-N,N-dibutylamino-6-methyl-7-anilino-fluorane

CVL: 3,3-bis(p-dimethylaminophenyl)-6-methyl-7-nitro-phthalide

5 NEW-Blue: 3-(4-diethylamino-2-ethoxyphenyl)-3-(1-ethyl-2-methylindol-3-yl)-4-azaphthalide

1-red: 3,3-bis(1-ethyl-2-methylindol-3-yl)phthalide

First, a dye precursor dispersion (Solution C) of the following composition was milled by a sand grinder to an average particle diameter of 1 micron.

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(Solution C: dye precursor dispersion)

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Above dye precursor	2.0 parts
10% Aqueous polyvinylalcohol solution	4.6
Water	2.6

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Then, the color developer dispersion used in Example 4 (or Example 6), Solution C, and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

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Color developer dispersion of Example 6 using the compound A8 (or color developer dispersion of Example 4 using the compound A6)	36.0 part
Solution C: dye precursor dispersion	9.2
Kaolin clay (50% dispersion)	12.0

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The coating color was coated on one side of a 50g/m² base paper in an amount of 6.0 g/m² and dried, and the sheet was supercalendered to a flatness of 500 to 600 seconds to obtain a thermal recording sheet.

Comparative Examples 1-7

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Thermal recording sheets for comparative tests were prepared using the known compounds shown below as color developers and using the same procedure as in Examples 1-48.

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(Table 4)

(Known color developer compounds)

Bisphenol A (B1)

Bisphenol S (B2)

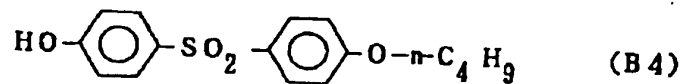
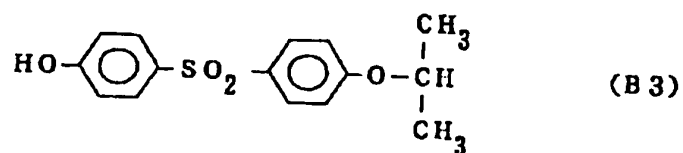
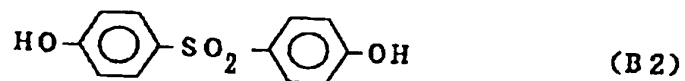
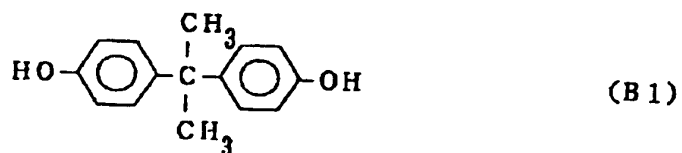
4-Hydroxy-4'-iso-propoxydiphenylsulfone (B3)

4-Hydroxy-4'-n-propoxydiphenylsulfone (B4)

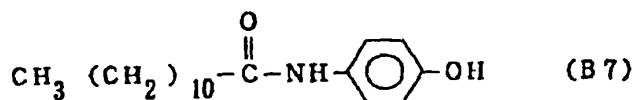
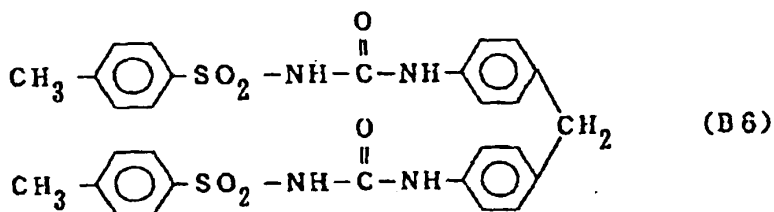
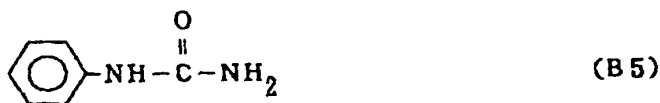
Phenylurea (B5) described in Japanese OPI 58-211496

Dimerized urea (B6) described in Japanese OPI 5-147357

Amidephenol derivative (B7)



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Specifically, dispersions of the above individual known color developer compounds of the following composition were milled by a sand grinder to an average particle diameter of 1 micron.

(Solution D: color developer dispersion)

Known color developer compound (B1-B7)	6.0 parts
10% Aqueous polyvinylalcohol solution	18.8
Water	11.2

Then, Solution D (color developer dispersion), the dye precursor (ODB) dispersion (Solution B) used in Examples 1-48, and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

Solution D: color developer dispersion	36.0 parts
Solution B: dye precursor dispersion (ODB dispersion)	9.2
Kaolin clay (50% dispersion)	12.0

The coating color was coated on one side of a 50g/m² base paper in an amount of 6.0 g/m² and dried, and the sheet was supercalendered to a flatness of 500 to 600 seconds to obtain a thermal recording sheet.

Comparative Examples 8-9

Thermal recording sheets for comparative tests were prepared using bisphenol A as a color developer, and ODB-2 and NEW-BLUE and using the same procedure as in Examples 49-56. (Table 5)

Evaluation of Thermal Recording Sheets

The resulting thermal recording sheets were tested for basic properties by a recordability test using a thermal printer, a ground color thermal stability test, and an oil ink adaptability test.

Recordability test (dynamic color developing density):

To test the recording adaptability, the thermal recording sheet was recorded using a word processor printer (RUPO-90F: Trade mark of word processor made by Toshiba co., ltd.) at a maximum energy, and the recorded portion was measured by a Macbeth densitometer (RD-914, an amber filter used. Hereinafter density was measured in this condition). In this case, the greater the Macbeth value, the higher the recording density and the better the recording adaptability.

Ground color thermal stability test (static color developing density):

To test the thermal stability of ground color, the recording sheet was pressed against a hot plate heated individually at 90°C, 120°C, and 150°C at a pressure of 10g/cm² for 5 seconds, and the treated sheet was measured by a Macbeth densitometer. In this case, the smaller the Macbeth value, the smaller the coloring of ground color and the higher the thermal stability of ground color.

Oil ink adaptability test (discoloration of ground color by oil ink):

The recording sheet was written with a felt pen of red oil ink (Magic INK No. 900/Teranishi Kagaku co., ltd.), and visually measured for a degree of discoloration compared to a conventional red ink.

- A: No discoloration
- B: Little discoloration
- C: Slight discoloration
- D: Considerable discoloration

The evaluation results of the basic properties of Examples 1-56 using the dimerized or trimerized urea compounds of the present invention as color developers, and Comparative Examples 1-9 using the conventional color developer compound are shown in Tables 1 to 5.

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Table 1

5	Entry	Color developer	Ground color before recording	Recordability test	Thermal stability of ground color			Oil ink adaptabil- ity
					90°C	120°C	150°C	
	Ex. 1	A3	0.03	1.34	0.03	0.03	0.04	A
	Ex. 2	A4	0.02	1.12	0.02	0.03	0.04	A
10	Ex. 3	A5	0.03	1.28	0.03	0.03	0.05	A
	Ex. 4	A6	0.02	1.26	0.02	0.03	0.04	A
	Ex. 5	A7	0.03	1.35	0.03	0.04	0.05	A
15	Ex. 6	A8	0.02	1.21	0.02	0.03	0.05	A
	Ex. 7	A10	0.04	1.15	0.04	0.04	0.04	A
	Ex. 8	A12	0.02	1.16	0.02	0.03	0.08	A
20	Ex. 9	A14	0.06	1.28	0.06	0.06	0.06	A
	Ex. 10	A15	0.02	1.16	0.03	0.03	0.05	A
	Ex. 11	A16	0.03	1.07	0.03	0.04	0.06	A
	Ex. 12	A17	0.02	1.29	0.02	0.04	0.24	A
25	Ex. 13	A19	0.03	1.31	0.03	0.05	0.74	A
	Ex. 14	A28	0.03	1.28	0.03	0.04	0.79	A
	Ex. 15	A31	0.03	1.15	0.03	0.04	0.05	A
30	Ex. 16	A32	0.03	1.24	0.03	0.04	0.05	A
Note: ODB used as a dye								

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Table 2

Entry	Color developer	Ground color before recording	Recordability test	Thermal stability of ground color			Oil ink adaptability
				90°C	120°C	150°C	
Ex.17	A33	0.04	0.92	0.04	0.05	0.06	A
Ex.18	A34	0.03	0.91	0.03	0.04	0.05	A
Ex.19	A49	0.04	1.22	0.04	0.04	0.05	A
Ex.20	A50	0.03	1.16	0.03	0.03	0.05	A
Ex.21	A51	0.03	0.98	0.03	0.04	0.05	A
Ex.22	A52	0.04	1.11	0.04	0.04	0.05	A
Ex.23	A53	0.03	1.10	0.03	0.03	0.05	A
Ex.24	A54	0.05	1.35	0.05	0.06	0.16	A
Ex.25	A55	0.03	1.23	0.03	0.04	0.05	A
Ex.26	A56	0.04	1.24	0.04	0.04	0.06	A
Ex.27	A63	0.03	0.82	0.03	0.04	0.05	A
Ex.28	A64	0.04	0.87	0.05	0.05	0.05	A
Ex.29	A65	0.10	0.83	0.10	0.12	0.02	A
Ex.30	A66	0.04	0.86	0.04	0.05	0.06	A
Ex.31	A68	0.03	0.81	0.03	0.04	0.05	A
Ex.32	A69	0.04	0.93	0.04	0.05	0.07	A
Note: ODB used s a dye							

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Table 3

Entry	Color developer	Ground color before recording	Recordability test	Thermal stability of ground color			Oil ink adaptability
				90°C	120°C	150°C	
Ex. 33	A79	0.05	0.85	0.05	0.06	0.12	A
Ex. 34	A82	0.12	0.80	0.12	0.13	0.14	A
Ex. 35	A90	0.03	0.85	0.03	0.04	0.05	A
Ex. 36	A92	0.03	0.85	0.03	0.03	0.03	A
Ex. 37	A100	0.02	1.14	0.02	0.04	0.05	A
Ex. 38	A101	0.03	1.09	0.03	0.04	0.05	A
Ex. 39	A103	0.02	0.90	0.03	0.04	0.06	A
Ex. 40	A104	0.02	0.92	0.02	0.04	0.28	A
Ex. 41	A111	0.03	1.00	0.03	0.04	0.05	A
Ex. 42	A112	0.04	0.81	0.04	0.04	0.05	A
Ex. 43	A120	0.05	0.82	0.05	0.05	0.08	A
Ex. 44	A124	0.02	0.74	0.03	0.04	0.05	A
Ex. 45	A186	0.04	1.07	0.04	0.04	0.04	A
Ex. 46	A200	0.02	1.16	0.02	0.03	0.05	A
Ex. 47	A201	0.03	1.03	0.03	0.04	0.05	A
Ex. 48	A203	0.04	1.03	0.05	0.07	0.09	A
Note: ODB used as a dye							

Table 4

Entry	Color developer	Ground color before recording	Recordability test	Thermal stability of ground color			Oil ink adaptability
				90°C	120°C	150°C	
Comp.Ex.1	B1	0.06	1.44	0.21	1.51	1.53	D
Comp.Ex.2	B2	0.06	1.30	0.08	0.21	0.58	D
Comp.Ex.3	B3	0.04	1.50	0.13	1.55	1.56	D
Comp.Ex.4	B4	0.04	1.52	0.04	0.13	1.55	D
Comp.Ex.5	B5	0.03	1.01	0.04	0.06	0.94	A
Comp.Ex.6	B6	0.06	1.14	0.07	0.11	1.02	A
Comp.Ex.7	B7	0.07	0.79	0.08	0.49	0.68	C
Note: ODB used as a dye							

Table 5

Entry	Dye	Color developer	Ground color before recording	Recordability test	Thermal stability of ground color			Oil ink adaptability
					90°C	120°C	150°C	
Ex.49	ODB-2	A8	0.05	1.26	0.05	0.05	0.07	A
Ex.50	CVL	A8	0.03	0.82	0.03	0.04	0.05	A
Ex.51	NEW-Blue	A8	0.07	0.98	0.07	0.08	0.10	A
Ex.52	I-red	A8	0.03	1.02	0.03	0.05	0.06	A
Ex.53	ODB-2	A6	0.05	1.30	0.05	0.06	0.07	A
Ex.54	CVL	A6	0.04	0.84	0.04	0.05	0.06	A
Ex.55	NEW-Blue	A6	0.08	0.90	0.08	0.09	0.09	A
Ex.56	I-red	A6	0.03	0.80	0.03	0.05	0.06	A
Com.Ex.8	ODB-2	B1	0.05	1.30	0.27	1.31	1.36	D
Com.Ex.9	NEW-Blue	B1	0.04	1.34	0.29	1.33	1.39	D

Then, to test for additional properties, the thermal recording sheet was subjected to a heat lamination test and a reversible recording test.

Heat lamination test (preparation of laminated recording sheet):

Using a simple lamination apparatus (MS Pouch H-140/ Meiko Shokai), the thermal recording sheet was put between pouch films to prepare a laminated thermal recording sheet, and the ground color thereof was measured by a Macbeth densitometer. In this case, the smaller the Macbeth value shows the better the ground color stability. In other words, the sheet can be laminated without coloring. The thermal recording sheet using the dimerized or trimerized urea compound of the present invention was able to be laminated with a stable ground color.

Reversible recording test:

As in the recordability test, the thermal recording sheet was recorded by a word processor printer, the recorded sheet was passed between 180°C heat rolls at a speed of 30mm/sec, and the recorded portion and the ground color portion were measured by a Macbeth densitometer. In this case, the smaller the Macbeth value of the recorded portion shows the higher the erasability. After that, the sheet was again recorded by the word processor printer, and the recorded portion was measured for Macbeth density.

Evaluation results of additional properties of the Examples using the dimerized or trimerized urea compound of the present invention as a color developer and Comparative Examples 1-8 using a known color developer compound are shown in Tables 6 to 9.

Table 6

Entry	Color developer	Heat lamination test	Reversible recordability test			Re-recording
			Recorded portion	Erasing (Recorded portion)	Ground color)	
Ex. 1	A3	0.10	1.34	0.20	0.04	1.31
Ex. 2	A4	0.11	1.12	0.13	0.04	1.09
Ex. 3	A5	0.12	1.28	0.18	0.04	1.24
Ex. 4	A6	0.11	1.26	0.16	0.04	1.24
Ex. 5	A7	0.12	1.35	0.19	0.05	1.27
Ex. 6	A8	0.11	1.21	0.14	0.04	1.19
Ex. 7	A10	0.12	1.15	0.14	0.04	1.10
Ex. 8	A12	0.11	1.16	0.32	0.16	1.11
Ex. 9	A14	0.14	1.28	0.13	0.06	1.20
Ex. 10	A15	0.10	1.16	0.37	0.04	1.11
Ex. 15	A31	0.12	1.15	0.20	0.04	1.10
Ex. 16	A32	0.12	1.24	0.19	0.04	1.18
Ex. 17	A33	0.13	0.92	0.22	0.06	0.88
Ex. 18	A34	0.10	0.91	0.13	0.04	0.86
Ex. 19	A49	0.14	1.22	0.19	0.04	1.19
Note: ODB used as a dye						

Table 7

Entry	Color developer	Heat lamination test	Reversible recordability test			Re-recording
			Recorded portion	Erasing (Recorded portion)	Ground color)	
Ex.20	A50	0.15	1.16	0.20	0.05	1.11
Ex.21	A51	0.12	0.96	0.40	0.05	0.93
Ex.22	A52	0.13	1.11	0.22	0.04	1.05
Ex.23	A53	0.14	1.10	0.21	0.05	1.02
Ex.25	A55	0.11	1.23	0.28	0.05	1.17
Ex.27	A63	0.13	0.82	0.50	0.06	0.78
Ex.28	A64	0.14	0.87	0.50	0.05	0.83
Ex.31	A68	0.12	0.81	0.44	0.05	0.79
Ex.33	A79	0.12	0.85	0.40	0.09	0.81
Ex.35	A90	0.12	0.79	0.35	0.05	0.72
Ex.37	A100	0.12	1.04	0.44	0.06	0.99
Ex.38	A101	0.12	1.09	0.43	0.04	1.01
Ex.43	A120	0.16	0.72	0.30	0.08	0.70
Note: ODB used as a dye						

Table 8

Entry	Color developer	Heat lamination	Reversible recordability test			Re-recording
			Recorded portion	Erasing (Recorded portion)	Ground color)	
Comp.Ex.1	B1	1.96	1.44	1.58	1.50	--
Comp.Ex.2	B2	0.37	1.30	1.43	0.93	--
Comp.Ex.3	B3	1.86	1.50	1.57	1.54	--
Comp.Ex.4	B4	0.28	1.52	1.54	1.49	--
Comp.Ex.5	B5	0.29	1.03	1.05	0.96	--
Comp.Ex.6	B6	0.28	1.14	1.11	0.99	--
Comp.Ex.7	B7	0.64	0.79	0.44	0.15	0.73
Note: ODB used as a dye						

Table 9

Entry	Dye	Color developer	Heat lamination	Revers. recordability test			Re-recording
				Recorded-Erasing	(Recorded portion	Ground color)	
Ex. 49	ODB-2	A8	0.09	1.26	0.10	0.05	1.25
Ex. 50	CVL	A8	0.10	0.82	0.11	0.05	0.80
Ex. 51	NEW-BLUE	A8	0.10	0.98	0.09	0.05	0.98
Ex. 52	I-red	A8	0.11	1.02	0.15	0.06	0.98
Comp.Ex.8	ODB-2	B1	1.84	1.30	1.56	1.48	- -
Comp.Ex.9	NEW-BLUE	B1	1.79	1.34	1.49	1.43	- -

Example 57

The thermal recording sheet of Example 1 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 1.20 and 0.07, respectively.

Example 58

The thermal recording sheet of Example 3 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 1.18 and 0.07, respectively.

Example 59

The thermal recording sheet of Example 4 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 1.04 and 0.08, respectively.

Example 60

The thermal recording sheet of Example 6 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 1.09 and 0.08, respectively.

Example 61

The thermal recording sheet of Example 49 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 1.12 and 0.07, respectively.

Example 62

The thermal recording sheet of Example 51 was subjected to 100 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 100th time were 0.94 and 0.09, respectively.

Example 63

The thermal recording sheet of Example 56 was subjected to 50 repetitions of a reversibility test by a heat roll. The Macbeth densities of the recorded portion and ground color of the 50th time were 0.76 and 0.11, respectively.

Example 64

The thermal recording sheet of Example 3 was recorded by a word processor printer, and the recorded surface was wiped out with ethanol. The recorded portion had a Macbeth density of 0.23.

Example 65

The thermal recording sheet of Example 9 was recorded by a word processor printer, and the recorded surface was wiped out with ethanol. The recorded portion had a Macbeth density of 0.16.

Example 66

The thermal recording sheet of Example 18 was recorded by a word processor printer, and the recorded surface was wiped out with ethanol. The recorded portion had a Macbeth density of 0.19.

Example 67

The thermal recording sheet of Example 3 was toner recorded by a copier (NP6060/ Canon co., ltd). No change in the ground color was noted.

Example 68

The thermal recording sheet of Example 5 was toner recorded by a copier (NP6060/ Canon co., ltd). No change in the ground color was noted.

Example 69

The thermal recording sheet of Example 28 was toner recorded by a copier (NP6060/ Canon co., ltd). No change in the ground color was noted.

Example 70

The thermal recording sheet of Example 53 was toner recorded by a copier (NP6060/ Canon co., ltd). No change in the ground color was noted. Production of Thermal recording sheet containing the optical absorbent.

Examples 71-77

As described above, an optical recording sheet was produced using 3-N,N-diethylamino-6-methyl-7-anilino-fluorane (ODB) as a dye precursor, the dimerized urea compound (or the trimerized urea compound) of the present invention as a color developer, and a heat melt (optical absorbent A) of a bis-dithiobenzylnickel complex and a sensitizer as an optical absorbent. (Table 10)

Specifically, 94 parts of 4-biphenyl-9-tolyether was mixed with 6 parts of bi-dithiobenzylnickel complex, heated to 100 to 150°C to melt, and then crushed to obtain an optical absorbent. An optical absorbent dispersion of the following composition was milled by a sand grinder to an average particle diameter of 1 micron.

(Solution E: optical absorbent dispersion)

Optical absorbent	4.0 parts
10% Aqueous polyvinylalcohol solution	10.0
Water	6.0.

Then, the color developer dispersion (Solution A) used in Examples 1-48, the dye precursor (ODB) dispersion (Solution B) used in Examples 1-48, Solution E (optical absorbent dispersion), and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

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Solution A (color developer dispersion)	36.0 parts
Solution B (dye precursor dispersion)	9.2
Solution E (optical absorbent dispersion)	20.0
Kaolin clay (50% dispersion)	12.0

The coating color was coated on one side of a 50g/m² base paper in an amount of 6.0 g/m² and dried, and the sheet was supercalendered to a flatness of 500 to 600 seconds to obtain an optical recording sheet.

Examples 78-81

An optical recording sheet was produced using NK-2612 (Nippon Kanko Shikiso Kenkyusho) (optical absorbent B) as an optical absorbent in place of the heat melt of bis-dithiobenzylnickel complex and a sensitizer. (Table 10)
First, the following aqueous optical absorbent solution was prepared.

(Solution F: aqueous optical absorbent solution)

NK-2612	0.04 parts
Water	3.96.

Then, the color developer dispersion (Solution A) used in Examples 1-48, the dye precursor (ODB) dispersion (Solution B) used in Examples 1-48, Solution E (optical absorbent dispersion), and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

Solution A (color developer dispersion)	36.0 parts
Solution B (dye precursor dispersion)	9.2
Solution F (optical absorbent solution)	4.0
Kaolin clay (50% dispersion)	12.0

The coating color was coated on one side of a 50g/m² base paper in an amount of 6.0 g/m² and dried, and the sheet was supercalendered to a flatness of 500 to 600 seconds to obtain an optical recording sheet.

Examples 82-84

An optical recording sheet was produced using toluenedithiolnickel complex (optical absorbent C) as an optical absorbent in place of the heat melt of bis-dithiobenzylnickel complex and a sensitizer.

First, an optical absorbent dispersion (Solution G) was milled by a sand grinder to an average particle diameter of 1 micron.

(Solution G: optical absorbent color developer dispersion)

Compound A6 (or compound A8 or A66)	6.0 parts
Toluenedithiolnickel complex	1.0
10% Aqueous polyvinylalcohol solution	18.8
Water	10.2.

Then, the optical absorbent color developer dispersion (Solution G) the dye precursor (ODB) dispersion (Solution

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B) used in Examples 1 -48, and a kaolin clay dispersion were mixed in the following ratio to obtain a coating color.

5

Solution G (optical absorbent color developer dispersion)	36.0 parts
Solution B (dye precursor dispersion)	9.2
Kaolin clay (50% dispersion)	12.0

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The coating color was coated on one side of a 50g/m² base paper in an amount of 6.0 g/m² and dried, and the sheet was supercalendered to a flatness of 500 to 600 seconds to obtain an optical recording sheet.

Examples 85-86

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Using ODB-2 or NEW-Blue as a dye precursor in place of ODB, the same procedure as in Examples 71-77 was used to obtain an optical recording sheet.

Examples 87-88

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Using ODB-2 or NEW-Blue as a dye precursor in place of ODB, the same procedure as in Examples 78-81 was used to obtain an optical recording sheet.

Examples 89-90

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Using NEW-Blue or I-red as a dye precursor in place of ODB, the same procedure as in Examples 82-84 was used to obtain an optical recording sheet.

Evaluation of optical recording sheets

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The resulting optical recording sheets were subjected to a recordability test.

Recordability test (optical recording):

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Using a laser plotter described in Japanese OPI 3-239598, the optical recording sheet was irradiated with a laser light, and the recorded portion was measured by a Macbeth densitometer. Using a 30 mW semiconductor laser LT015MD (Sharp) with an oscillation wavelength of 830 nm as a recording light sources, an aspheric plastic lens AP4545 (Konica) with a numerical aperture of 0.45 and a focal length of 4.5 mm as an optical converging lens, a recording speed of 50 mm/sec, and a recording interval of 50 microns, a 1 cm square overall recording was obtained. The evaluation results are shown in Table 10.

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Table 10

Entry	Color developer	Dye	Optical absorbent	Recordability test
Example 71	Compound A3	ODB	Absorbent A	1.38
Example 72	Compound A6	ODB	Absorbent A	1.28
Example 73	Compound A8	ODB	Absorbent A	1.25
Example 74	Compound A19	ODB	Absorbent A	1.40
Example 75	Compound A39	ODB	Absorbent A	1.00
Example 76	Compound A49	ODB	Absorbent A	1.22
Example 77	Compound A68	ODB	Absorbent A	1.02
Example 78	Compound A6	ODB	Absorbent B	1.26
Example 79	Compound A7	ODB	Absorbent B	1.33
Example 80	Compound A16	ODB	Absorbent B	1.10
Example 81	Compound A63	ODB	Absorbent B	1.04
Example 82	Compound A6	ODB	Absorbent C	1.27
Example 83	Compound A8	ODB	Absorbent C	1.26
Example 84	Compound A66	ODB	Absorbent C	1.01
Example 85	Compound A6	ODB	Absorbent A	1.27
Example 86	Compound A6	ODB	Absorbent A	1.02
Example 87	Compound A6	ODB	Absorbent B	1.26
Example 88	Compound A6	ODB	Absorbent B	1.00
Example 89	Compound A6	ODB	Absorbent C	1.01
Example 90	Compound A8	ODB	Absorbent C	0.98
Absorbent A: bis-dithiobenzylnickel complex/sensitizer				
Absorbent B: NK-2612				
Absorbent C: toluenedithiolnickel complex				

As described above, it can be seen that the dimerized or trimerized urea compound of the present invention is an epoch-making color developer that can provide a recording with a practical image density by a thermal head or the like while without a substantial change in ground color at environmental temperatures of 120 to 150°C. Therefore, the present invention has the following effects.

- (1) A thermal recording sheet superior in storage stability such as heat resistance and solvent resistance compared to conventional thermal recording sheets.
- (2) The thermal recording sheet can be used under severe conditions (e.g. at temperatures of 90-150°C) at which conventional products could not be used.
- (3) Since the thermal recording sheet does not undergo discoloration when written with an oil ink, it can be freely written using these writing means.
- (4) The thermal recording sheet can be simply heat laminated by a simple laminator or the like. Cards and the like can also be easily prepared.
- (5) The thermal recording sheet can be toner recorded since the ground color is stable even when passed through a heat roll.

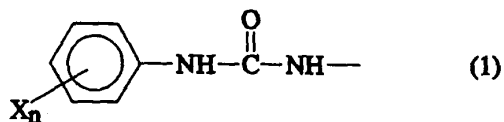
Further, the thermal recording sheet having an erasing function has the following advantages:

- (6) A new recording system, which enables repeated color recording and erasing, which leads to resource-saving.
- (7) Some types of the sheet can be erased merely by passing through a heat roll, without strict temperature control.
- (8) Unlike liquid crystals, the thermal recording sheet can be used a simple display means that can be recorded and erased using different thermal energies.

The thermal recording sheet of the present invention can also be incorporated with an optical absorbent for use as an optical recording sheet having similar effects.

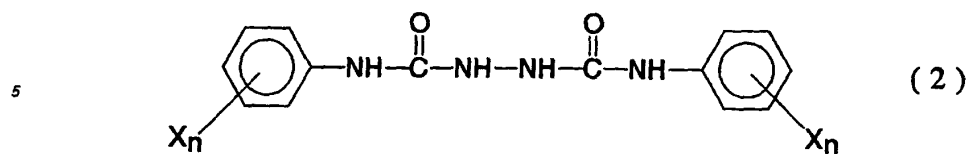
Claims

1. A thermal recording sheet which comprises a thermal recording layer comprising a colourless or pale coloured dye precursor and a colour developer, which colour developer is a urea compound which comprises at least two groups of Formula (I):

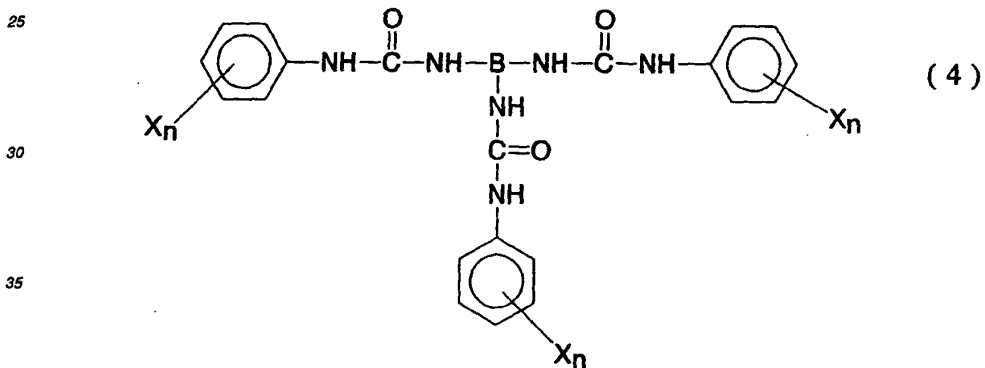
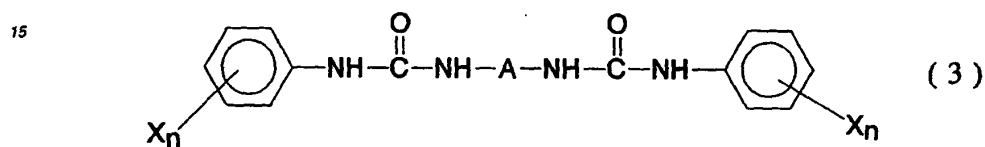


wherein X is C₁-C₁₂ alkyl, halogenated C₁-C₆ alkyl, C₇-C₁₄ aralkyl, C₁-C₁₂ alkoxy, C₆-C₁₂ aryloxy, C₂-C₁₂ alkoxy-carbonyl, C₁-C₁₂ acyl, C₂-C₁₂ dialkylamino, C₇-C₁₂ arylalkylamino, C₆-C₁₂ arylamino, C₁-C₁₂ acylamino, nitro, cyano, a halogen or hydrogen, and n is an integer of 1 to 3.

2. A thermal recording sheet according to claim 1 wherein the colour developer is a urea compound of Formula (2), (3) or (4)

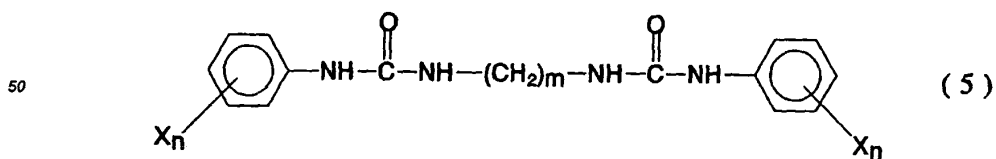


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wherein X is as defined in claim 1, A is a divalent group comprising 30 or less carbon atoms, B is a trivalent group comprising 30 or less carbon atoms and n is an integer from 1 to 3.

- 45 3. A thermal recording sheet according to claim 1 wherein the color developer is a urea compound of Formula (5)



wherein X is as defined in claim 1, n is an integer from 1 to 3, and m is an integer from 1 to 12.

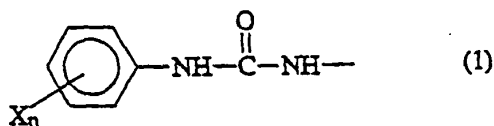
4. A thermal recording sheet according to claim 2 wherein the divalent group (A) of the color developer of Formula (3)

comprises:

- (a) a branched chain alkylene group having from 1 to 15 carbon atoms;
 - (b) two or more alkylene groups having from 1 to 12 carbon atoms, the alkylene groups being linked to each other via nitrogen or oxygen atoms;
 - (c) a group containing at least one substituted or unsubstituted cycloalkyl group having from 3 to 20 carbon atoms;
 - (d) a group having one substituted or unsubstituted aromatic ring having from 6 to 20 carbon atoms;
 - (e) a group having at least two aromatic rings, each of which may be substituted or unsubstituted, which group has from 12 to 30 carbon atoms; or
 - (f) a group containing a substituted or unsubstituted piperazine ring and having from 4 to 12 carbon atoms.
5. An optical recording sheet which comprises a thermal recording sheet according to any one of claims 1 to 4 and an optical absorbent
 6. A thermal recording card which comprises a thermal recording sheet according to any one of claims 1 to 4 or an optical recording sheet according to claim 5, the said thermal or optical sheet being laminated with a plastic film.
 7. An electrophotographic transfer sheet which comprises a thermal recording sheet according to any one of claims 1 to 4 or an optical recording sheet according to claim 5.
 8. A method of reversibly recording an image on a substrate, which method comprises
 - recording an image on a thermal recording sheet as defined in any one of claims 1 to 4 or an optical recording sheet as defined in claim 5;
 - erasing the recorded image by applying respectively heat or light to the recorded portion;
 - and recording an image again on the said sheet.
 9. A method of reversibly recording an image on a substrate, which method comprises
 - recording an image on a thermal recording sheet as defined in any one of claims 1 to 4 or an optical recording sheet as defined in claim 5;
 - contacting the recorded portion with an alcoholic solvent to erase the recorded image;
 - and recording an image again on the said sheet.

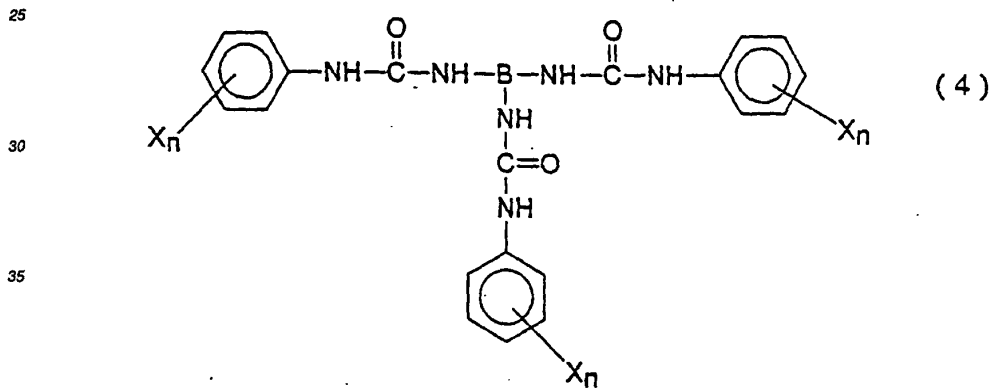
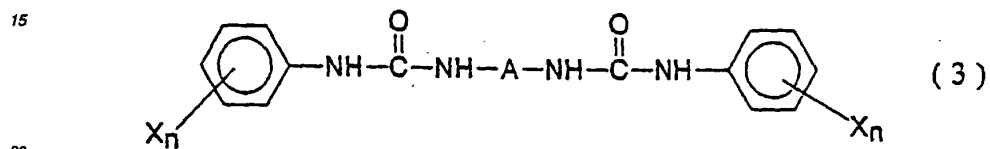
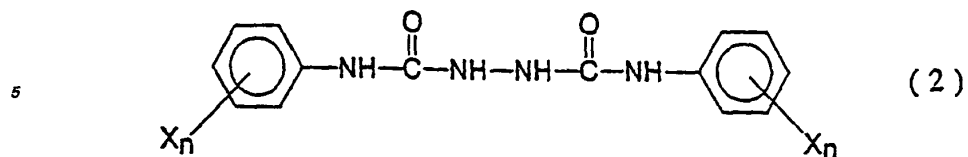
Patentansprüche

1. Wärmeempfindliches Aufzeichnungsblatt, das eine wärmeempfindliche Aufzeichnungsschicht umfaßt, die einen farblosen oder schwach gefärbten Farbstoffvorläufer und einen Farbentwickler umfaßt, wobei der Farbentwickler eine Harnstoffverbindung ist, die mindestens zwei Gruppen der Formel (1) umfaßt:



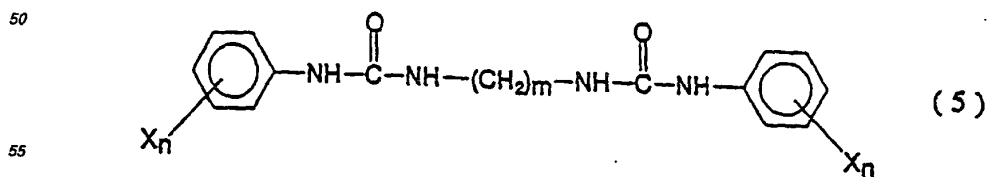
worin X C₁-C₁₂-Alkyl, halogeniertes C₁-C₆-Alkyl, C₇-C₁₄-Aralkyl, C₁-C₁₂-Alkoxy, C₆-C₁₂-Aryloxy, C₂-C₁₂-Alkoxy-carbonyl, C₁-C₁₂-Acyl, C₂-C₁₂-Dialkylamino, C₇-C₁₂-Arylalkylamino, C₆-C₁₂-Arylamino, C₁-C₁₂-Acylamino, Nitro, Cyan, ein Halogen oder Wasserstoff ist, und n eine ganze Zahl von 1 bis 3 bedeutet.

2. Wärmeempfindliches Aufzeichnungsblatt nach Anspruch 1, dadurch gekennzeichnet, daß der Farbentwickler eine Harnstoffverbindung der Formel (2), (3) oder (4) ist



45 worin X die im Anspruch 1 angegebene Bedeutung besitzt, A eine zweiwertige Gruppe mit 30 oder weniger Kohlenstoffatomen ist, B eine dreiwertige Gruppe mit 30 oder weniger Kohlenstoffatomen ist und n eine ganze Zahl von 1 bis 3 bedeutet.

3. Wärmeempfindliches Aufzeichnungsblatt nach Anspruch 1, dadurch gekennzeichnet, daß der Farbtentwickler eine Harnstoffverbindung der Formel (5) ist



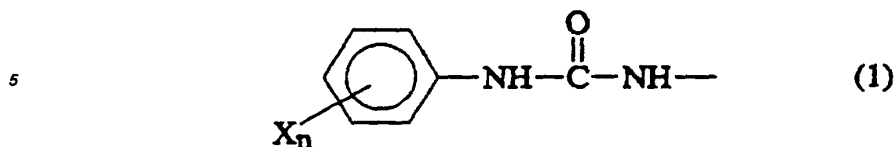
worin X die in Anspruch 1 angegebene Bedeutung besitzt, n eine ganze Zahl von 1 bis 3 ist, und m eine ganze Zahl von 1 bis 12 ist.

- 5 4. Wärmeempfindliches Aufzeichnungsblatt nach Anspruch 2, dadurch gekennzeichnet, daß die zweiwertige Gruppe (A) des Farhentwicklers der Formel (3) umfaßt:
 - (a) eine verzweigte Alkylengruppe mit 1 bis 15 Kohlenstoffatomen;
 - (b) zwei oder mehrere Alkylengruppen mit 1 bis 12 Kohlenstoffatomen, wobei die Alkylengruppen miteinander über Stickstoff- oder Sauerstoffatome gebunden sind;
 - 10 (c) eine Gruppe, die mindestens eine substituierte oder unsubstituierte Cycloalkylgruppe mit 3 bis 20 Kohlenstoffatomen enthält;
 - (d) eine Gruppe, die einen substituierten oder unsubstituierten aromatischen Ring mit 6 bis 20 Kohlenstoffatomen besitzt;
 - 15 (e) eine Gruppe, die mindestens zwei aromatische Ringe besitzt, von denen jeder substituiert oder unsubstituiert sein kann, wobei die Gruppe 12 bis 30 Kohlenstoffatome besitzt; oder
 - (f) eine Gruppe, die einen substituierten oder unsubstituierten Piperazinring enthält und 4 bis 12 Kohlenstoffatome besitzt.
- 20 5. Optisches Aufzeichnungsblatt, das ein wärmeempfindliches Aufzeichnungsblatt nach einem der Ansprüche 1 bis 4 und einen optischen Absorber umfaßt.
6. Wärmeempfindliche Aufzeichnungskarte, die ein wärmeempfindliches Aufzeichnungsblatt nach einem der Ansprüche 1 bis 4 oder ein optisches Aufzeichnungsblatt nach Anspruch 5 umfaßt, wobei das wärmeempfindliche oder optische Blatt mit einem Kunststoffilm laminiert ist.
- 25 7. Elektrophotographisches Übertragungsblatt, das ein wärmeempfindliches Aufzeichnungsblatt gemäß einem der Ansprüche 1 bis 4 oder ein optisches Aufzeichnungsblatt nach Anspruch 5 umfaßt.
8. Verfahren zur reversiblen Aufzeichnung eines Bildes auf einem Substrat, dadurch gekennzeichnet, daß das Verfahren umfaßt:
 - Aufzeichnen eines Bildes auf einem wärmeempfindlichen Aufzeichnungsblatt nach einem der Ansprüche 1 bis 4 oder einem optischen Aufzeichnungsblatt nach Anspruch 5;
 - 35 Auslöschen des aufgezeichneten Bildes durch Applikation von Hitze oder Licht auf dem mit der Aufzeichnung versehenen Teil;
 - und wiederholte Aufzeichnung eines Bildes auf dem Blatt.
9. Verfahren zur reversiblen Aufzeichnung eines Bildes auf einem Substrat, dadurch gekennzeichnet, daß das Verfahren umfaßt:
 - 40 Aufzeichnen eines Bildes auf einem thermischen Aufzeichnungsblatt nach einem der Ansprüche 1 bis 4 oder einem optischen Aufzeichnungsblatt nach Anspruch 5;
 - In-Kontakt-Bringen des mit der Aufzeichnung versehenen Teils mit einem alkoholischen Lösungsmittel, um das aufgezeichnete Bild auszulöschen;
 - 45 und wiederholte Aufzeichnung eines Bildes auf dem Blatt.

Revendications

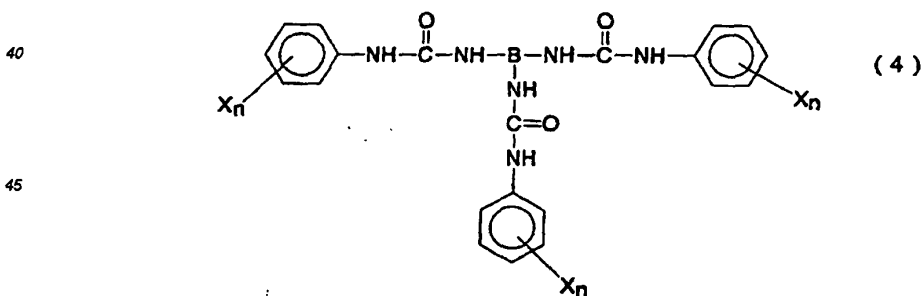
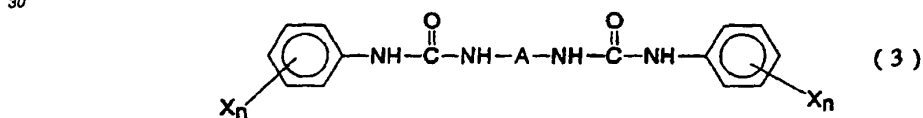
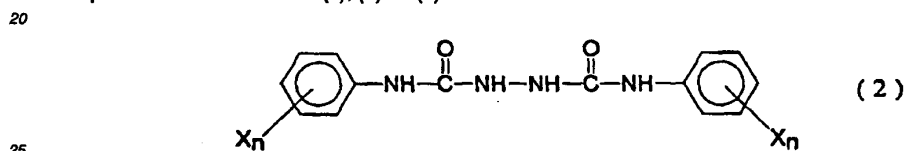
- 50 1. Feuille d'enregistrement thermique qui comprend une couche d'enregistrement thermique comprenant un précurseur de colorant incolore ou de couleur pâle et un développeur de couleur, lequel développeur de couleur est un composé d'urée qui comprend au moins deux groupes de la formule (1) :

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10 dans laquelle X est un alcoyle à 1 à 12 atomes de carbone, un alcoyle à 1 à 6 atomes de carbone halogéné, un aralcoyle à 7 à 14 atomes de carbone, un alcoxy à 1 à 12 atomes de carbone, un aryloxy à 6 à 12 atomes de carbone, un alcoxy-carbonyl à 2 à 12 atomes de carbone, un acyle à 1 à 12 atomes de carbone, un dialcoylamino à 2 à 12 atomes de carbone, un arylalcoylamino à 7 à 12 atomes de carbone, un arylamino à 6 à 12 atomes de carbone, un acylamino à 1 à 12 atomes de carbone, un nitro, un cyano, un halogène ou un hydrogène, et n est un entier de 1 à 3.

2. Feuille d'enregistrement thermique selon la revendication 1, dans laquelle le développeur de couleur est un composé défini par la formule (2), (3) ou (4) :

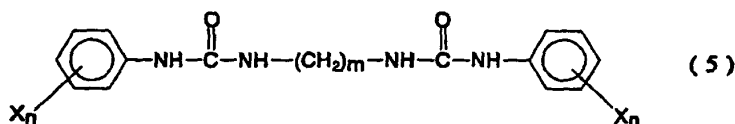


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dans laquelle X est comme défini dans la revendication 1, A est un groupe divalent comprenant 30 atomes de carbone ou moins, B est un groupe trivalent comprenant 30 atomes de carbone ou moins et n est un entier de 1 à 3.

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3. Feuille d'enregistrement thermique selon la revendication 1, dans laquelle le développeur de couleur est un composé défini par la formule (5) :



10 dans laquelle X est comme défini dans la revendication 1, n est un entier de 1 à 3 et m est un entier de 1 à 12.

- 10 4. Feuille d'enregistrement thermique selon la revendication 2, dans laquelle le groupe divalent (A) du développeur de couleur de la formule (3) comprend :

- 15 (a) un groupe alcoylène à chaîne ramifiée ayant 1 à 15 atomes de carbone ;
 (b) deux groupes alcoylènes ou davantage ayant 1 à 12 atomes de carbone, les groupes alcoylènes étant liés les uns aux autres par des atomes d'azote ou d'oxygène ;
 (c) un groupe contenant au moins un groupe cycloalcoyle, substitué ou non substitué, ayant 3 à 20 atomes de carbone ;
 20 (d) un groupe ayant un cycle aromatique, substitué ou non substitué, ayant 6 à 20 atomes de carbone ;
 (e) un groupe ayant au moins deux cycles aromatiques, dont chacun peut être substitué ou non substitué, lequel groupe a 12 à 30 atomes de carbone ; ou
 (f) un groupe contenant un cycle pipérazine, substitué ou non substitué, ayant 4 à 12 atomes de carbone.

- 25 5. Feuille d'enregistrement optique comprenant une feuille d'enregistrement thermique selon l'une quelconque des revendications 1 à 4 et un absorbant optique.

- 30 6. Carte d'enregistrement thermique comprenant une feuille d'enregistrement thermique selon l'une quelconque des revendications 1 à 4 ou une feuille d'enregistrement optique selon la revendication 5, ladite feuille thermique ou optique étant stratifiée sur un film plastique.

7. Feuille de transfert électrophotographique comprenant une feuille d'enregistrement thermique selon l'une quelconque des revendications 1 à 4 ou une feuille d'enregistrement optique selon la revendication 5.

- 35 8. Procédé pour enregistrer de façon réversible une image sur un substrat, lequel procédé comprend les étapes suivantes :

- 40 - enregistrer une image sur une feuille d'enregistrement thermique comme définie dans l'une quelconque des revendications 1 à 4 ou sur une feuille d'enregistrement optique comme définie dans la revendication 5 ;
 - effacer l'image enregistrée par application, respectivement, de chaleur ou de lumière sur la portion enregistrée ; et
 - enregistrer à nouveau une image sur ladite feuille.

- 45 9. Procédé pour enregistrer de façon réversible une image sur un substrat, lequel procédé comprend les étapes suivantes :

- 50 - enregistrer une image sur une feuille d'enregistrement thermique comme définie dans l'une quelconque des revendications 1 à 4 ou sur une feuille d'enregistrement optique comme définie dans la revendication 5 ;
 - mettre en contact la portion enregistrée avec un solvant alcoolique pour effacer l'image enregistrée ; et
 - enregistrer à nouveau une image sur ladite feuille.